

SUBSTITUTED PYRAZOLES

Field of the Invention

5 This invention relates to a series of substituted pyrazoles, pharmaceutical compositions containing these compounds, and intermediates used in their manufacture, and methods of using them.

Background of the Invention

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Cathepsin S (EC 3.4.22.27) is a cysteine protease of the papain family found primarily in lysosomes (Bromme, D.; McGrath, M. E. High Level Expression and Crystallization of Recombinant Human Cathepsin S. *Protein Science* **1996**, 5, 789-791).

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The role of cathepsin S in the immune response is anticipated by its tissue distribution: cathepsin S is found primarily in lymphatic tissues, lymph nodes, the spleen, B lymphocytes, and macrophages (Kirschke, H. Chapter 211. Cathepsin S. In Handbook of Proteolytic Enzymes. Barrett, A. J.; Rawlings, N. D.; Woessner, J. F., Eds. San Diego: Academic Press, 1998. pp. 621-624.). Cathepsin S inhibitors have been shown in animal models to modulate antigen presentation and are effective in an animal model of asthma (Riese, R. J.; Mitchell, R. N.; Villadangos, J. A.; Shi, G.-P.; Palmer, J. T.; Karp, E. R.; De Sanctis, G. T.; Ploegh, H. L.; Chapman, H. A. Cathepsin S Activity
20 Regulates Antigen Presentation and Immunity. *J. Clin. Invest.* **1998**, 101, 2351-2363 and Shi, G.-P.; Villadangos, J. A.; Dranoff, G.; Small, C.; Gu, L.; Haley, K. J.; Riese, R.; Ploegh, H. L.; Chapman, H. A. Cathepsin S Required for Normal MHC Class II Peptide Loading and Germinal Center Development. *Immunity* **1999**, 10, 197-206.).

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Mice in which the gene encoding cathepsin S has been knocked out are less susceptible to collagen-induced arthritis and their immune systems have

an impaired ability to respond to antigens (Nakagawa, T. Y.; Brissette, W. H.; Lira, P. D.; Griffiths, R. J.; Petrushova, N.; Stock, J.; McNeish, J. D.; Eastman, S. E.; Howard, E. D.; Clarke, S. R. M.; Rosloniec, E. F.; Elliott, E. A.; Rudensky, A. Y. Impaired Invariant Chain Degradation and Antigen Presentation and Diminished Collagen-Induced Arthritis in Cathepsin S Null Mice. *Immunity* **1999**, 10, 207-217).

These data demonstrate that compounds that inhibit the proteolytic activity of human cathepsin S should find utility in the treatment of chronic autoimmune diseases including, but not limited to, lupus, rheumatoid arthritis, and asthma; and have potential utility in modulating the immune response to tissue transplantation.

There are a number of cathepsin S inhibitors reported in the literature. The most important patents are listed below.

Certain dipeptidyl nitriles are claimed by Novartis as cathepsin S inhibitors in: Altmann, et. al. WO-99/24460.

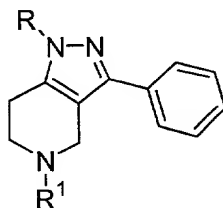
Dipeptidyl vinyl sulfones are claimed by Arris (now Axys) as cysteine protease (including cathepsin S) inhibitors in: Palmer, et. al. US-5976858.

Certain peptidyl sulfonamides are claimed by Arris/Axys as cysteine protease (including cathepsin S) inhibitors in: Palmer, et. al. US-5776718 (assigned to Arris, now Axys) & Klaus, et. al. US-6030946 (assigned to Axys).

Compounds somewhat similar to those of the present invention are described in the following references.

Winters, et. al. (Winters, G.; Sala, A.; Barone, D.; Baldoli, E. *J. Med. Chem.* **1985**, 28, 934-940; Singh, P.; Sharma, R. C. *Quant. Struct.-Act. Relat.* **1990**, 9, 29-32; Winters, G.; Sala, A.; Barone, D. in US-4500525 (1985)) have described bicyclic pyrazoles of the type shown below. R never contains a

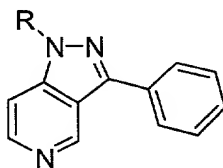
heterocyclic ring and no protease inhibitor activity is ascribed to these molecules; they are described as α 1-adrenergic receptor modulators.



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Shutske, et. al. claim the bicyclic pyrazoles below. The pyridine ring is aromatic in their system (Shutske, G. M.; Kapples, K. J.; Tomer, J. D. US-5264576 (1993)). Although reference is made to R being a linker to a heterocycle, the claims specify only R = hydrogen. The compounds are referred to as serotonin reuptake inhibitors.

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The compound 2-[4-[4-(3-methyl-5-phenyl-1H-pyrazol-1-yl)butyl]-1-piperazinyl]-pyrimidine is known from EP-382637, which describes pyrimidines having anxiolytic properties. This compound and analogs are further described in EP-502786 as cardiovascular and central nervous system agents. Pharmaceutical formulations with such compounds are disclosed in EP-655248 for use in the treatment of gastric secretion and as anti-ulcer agents. WO-9721439 describes medicaments with such compounds for treating obsessive-compulsive disorders, sleep apnea, sexual dysfunctions, emesis and motion sickness.

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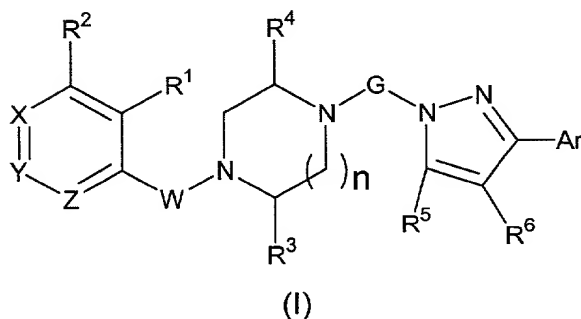
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The compounds 5-methyl-3-phenyl-1-[4-(4-phenyl-1-piperazinyl)butyl]-1H-indazole and 5-bromo-3-(2-chlorophenyl)-1-[4-(4-phenyl-1-piperazinyl)butyl]-1H-indazole, in particular the hydrochloride salts thereof, are known from WO-9853940 and CA 122:314528, where these and similar

compounds are described as kinase inhibitors in the former reference and possessing affinity for benzodiazepine receptors in the latter reference.

Summary of the Invention

The present invention concerns compounds which can be represented
5 by formula (I):



wherein:

- 10 R^1 is hydrogen, azido, halogen, C_{1-5} alkoxy, hydroxy, C_{1-5} alkyl, C_{2-5} alkenyl, cyano, nitro, R^7R^8N , C_{2-8} acyl, $R^9OC=O$, $R^{10}R^{11}NC=O$, or $R^{10}R^{11}NSO_2$; or R^1 is taken together with W as described below;
- R^2 is hydrogen, halogen, C_{1-5} alkoxy, C_{1-5} alkyl, C_{2-5} alkenyl, C_{1-5} haloalkyl, cyano, or $R^{48}R^{49}N$;
- 15 alternatively, R^1 and R^2 can be taken together to form an optionally substituted 5- to 7- membered carbocyclic or heterocyclic ring, which ring may be unsaturated or aromatic;
- each of R^3 and R^4 is independently hydrogen or C_{1-5} alkyl;
- each of R^5 and R^6 is independently hydrogen, C_{1-5} alkyl, C_{2-5} alkenyl, C_{1-5} alkoxy, C_{1-5} alkylthio, halogen, or a 4-7 membered carbocyclyl or
- 20 heterocyclyl;
- alternatively, R^5 and R^6 can be taken together to form an optionally substituted 5- to 7- membered carbocyclic or heterocyclic ring, which ring may be unsaturated or aromatic, and may be optionally substituted with between
- 25 one and three substituents independently selected from halo, cyano, amino, nitro, R^{40} , $R^{40}O-$, $R^{40}S-$, $R^{40}O(C_{1-5} \text{ alkylene})-$, $R^{40}O(C=O)-$, $R^{40}(C=O)-$, $R^{40}(C=S)-$, $R^{40}(C=O)O-$, $R^{40}O(C=O)(C=O)-$, $R^{40}SO_2$, $NHR^{62}(C=NH)-$, $NHR^{62}SO_2-$, and $NHR^{62}(C=O)-$;

R⁴⁰ is H, C₁₋₅ alkyl, C₂₋₅ alkenyl, phenyl, benzyl, phenethyl, C₁₋₅ heterocyclyl, (C₁₋₅ heterocyclyl)C₁₋₅ alkylene, amino, or mono- or di(C₁₋₅ alkyl)amino, or R⁵⁸OR⁵⁹-, wherein R⁵⁸ is H, C₁₋₅ alkyl, C₂₋₅ alkenyl, phenyl, benzyl, phenethyl, C₁₋₅ heterocyclyl, or (C₁₋₅ heterocyclyl)C₁₋₆ alkylene and R⁵⁹

is C₁₋₅ alkylene, phenylene, or divalent C₁₋₅ heterocyclyl; and

R⁶² can be H in addition to the values for R⁴⁰;

R⁷ is hydrogen, C₁₋₅ alkyl, C₃₋₅ alkenyl, phenyl, naphthyl, C₁₋₅ heterocyclyl, C₂₋₈ acyl, aroyl, R²⁷OC=O, R²⁸R²⁹NC=O, R²⁷SO, R²⁷SO₂, or R²⁸R²⁹NSO₂;

R⁸ is hydrogen, C₁₋₅ alkyl, C₃₋₅ alkenyl, phenyl, or C₁₋₅ heterocyclyl;

alternatively, R⁷ and R⁸ can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring may be saturated, unsaturated or aromatic;

R⁹ is C₁₋₅ alkyl, phenyl, naphthyl, or C₁₋₅ heterocyclyl;

R²¹ is hydrogen, C₁₋₅ alkyl, C₃₋₅ alkenyl, phenyl, naphthyl, C₁₋₅ heterocyclyl, C₂₋₈ acyl, aroyl, R³⁰OC=O, R³¹R³²NC=O, R³⁰SO, R³⁰SO₂, or R³¹R³²NSO₂;

R²² is hydrogen, C₁₋₅ alkyl, C₃₋₅ alkenyl, phenyl, or C₁₋₅ heterocyclyl;

alternatively, R²¹ and R²² can be taken together to form an optionally substituted 4- to 7-membered heterocyclic ring, which ring may be saturated, unsaturated or aromatic;

each of R²³, R²⁶, R²⁷, R³⁰, R³³, R⁴⁴, R⁴⁵, and R⁵⁰ is C₁₋₅ alkyl, phenyl, naphthyl, or C₁₋₅ heterocyclyl;

R²⁴ is hydrogen, C₁₋₅ alkyl, C₃₋₅ alkenyl, phenyl, naphthyl, C₁₋₅ heterocyclyl, C₂₋₈ acyl, aroyl, R³³OC=O, R³⁴R³⁵NC=O, R³³SO, R³³SO₂, or R³⁴R³⁵NSO₂;

R²⁵ is hydrogen, C₁₋₅ alkyl, C₃₋₅ alkenyl, phenyl, or C₁₋₅ heterocyclyl;

alternatively, R²⁴ and R²⁵ can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring may be saturated, unsaturated or aromatic;

each of R¹⁰ and R¹¹ is independently hydrogen, C₁₋₅ alkyl, C₂₋₅ alkenyl, phenyl, or C₁₋₅ heterocyclyl;

alternatively, R¹⁰ and R¹¹ or can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring may be saturated, unsaturated or aromatic;

each of R²⁸, R²⁹, R³¹, R³², R³⁴, R³⁵, R⁴⁶, R⁴⁷, R⁵¹ and R⁵² is independently

hydrogen, C₁₋₅ alkyl, phenyl, or C₁₋₅ heterocyclyl;

alternatively, R²⁸ and R²⁹, R³¹ and R³², R³⁴ and R³⁵, R⁴⁶ and R⁴⁷, or R⁵¹ and R⁵², independently, can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring may be

n is 1 or 2;

G represents C₃₋₆ alkenediyl or C₃₋₆ alkanediyl, optionally substituted with hydroxy, halogen, C₁₋₅ alkyl, C₁₋₅ alkoxy, oxo, hydroximino, CO₂R⁶⁰, R⁶⁰R⁶¹NCO₂, (L)-C₁₋₄ alkylene-, (L)-C₁₋₅ alkoxy, N₃, or [(L)-C₁₋₅ alkylene]amino;

each of R⁶⁰ and R⁶¹ is independently hydrogen, C₁₋₅ alkyl, C₃₋₅ alkenyl, phenyl, benzyl, phenethyl, or C₁₋₅ heterocyclyl; alternatively R⁶⁰ and R⁶¹, can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring may be saturated, unsaturated or aromatic;

L is amino, mono- or di-C₁₋₅ alkylamino, pyrrolidinyl, morpholinyl, piperidinyl homopiperidinyl, or piperazinyl, where available ring nitrogens may be optionally substituted with C₁₋₅ alkyl, benzyl, C₂₋₅ acyl, C₁₋₅ alkylsulfonyl or C₁₋₅ alkyloxycarbonyl;

X is nitrogen or R¹²C;

Y is nitrogen or R¹³C;

Z is nitrogen or R¹⁴C;

R¹² is hydrogen, halogen, C₁₋₅ alkoxy, C₁₋₅ alkyl, C₂₋₅ alkenyl, cyano, nitro, R²¹R²²N, C₂₋₈ acyl, C₁₋₅ haloalkyl, C₁₋₅ heterocyclyl, (C₁₋₅ heterocyclyl)C₁₋₅ alkylene, R²³OC=O, R²³O(C=O)NH-, R²³SO, R²²NHCO-, R²²NH(C=O)NH-, R²³(C₁₋₄ alkylene)NHCO-, R²³SO₂, or R²³SO₂NH-;

R¹³ is hydrogen, halogen, C₁₋₅ alkoxy, C₁₋₅ alkyl, C₂₋₅ alkenyl, cyano, nitro, R⁴²R⁴³N, C₂₋₈ acyl, C₁₋₅ haloalkyl, C₁₋₅ heterocyclyl, (C₁₋₅ heterocyclyl)C₁₋₅ alkylene, R⁴⁴OC=O, R⁴⁴O(C=O)NH-, R⁴⁴SO, R⁴³NHCO-, R⁴³NH(C=O)NH-, R⁴⁴(C₁₋₄ alkylene)NHCO-, R⁴⁴SO₂, or R⁴⁴SO₂NH-;

R¹⁴ is hydrogen, halogen, C₁₋₅ alkoxy, C₁₋₅ alkyl, C₂₋₅ alkenyl, cyano, nitro, R²⁴R²⁵N, C₂₋₈ acyl, C₁₋₅ haloalkyl, C₁₋₅ heterocyclyl, (C₁₋₅ heterocyclyl)C₁₋₅ alkylene, R²⁶OC=O, R²⁶O(C=O)NH-, R²⁶SO, R²⁵NHCO-, R²⁵NH(C=O)NH-, R²⁶(C₁₋₄ alkylene)NHCO-, R²⁶SO₂, or R²⁶SO₂NH-;

alternatively, R^{12} and R^{13} or R^{12} and R^2 or R^{13} and R^{14} can be taken together to form an optionally substituted 5- to 6- membered carbocyclic or heterocyclic ring, which ring may be unsaturated or aromatic;

Ar represents a monocyclic or bicyclic aryl or heteroaryl ring, optionally substituted with between 1 and 3 substituents selected from halogen, C_{1-5} alkoxy, C_{1-5} alkyl, C_{2-5} alkenyl, cyano, azido, nitro, $R^{15}R^{16}N$, $R^{17}SO_2$, $R^{17}S$, $R^{17}SO$, $R^{17}OC=O$, $R^{15}R^{16}NC=O$, C_{1-5} haloalkyl, C_{1-5} haloalkoxy, C_{1-5} haloalkylthio, and C_{1-5} alkylthio;

R^{15} is hydrogen, C_{1-5} alkyl, C_{3-5} alkenyl, phenyl, benzyl, C_{1-5} heterocyclyl, C_{2-8} acyl, aroyl, $R^{53}OC=O$, $R^{54}R^{55}NC=O$, $R^{53}S$, $R^{53}SO$, $R^{53}SO_2$, or $R^{54}R^{55}NSO_2$;

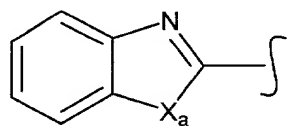
R^{16} is hydrogen, C_{1-5} alkyl, C_{3-5} alkenyl, phenyl, benzyl, or C_{1-5} heterocyclyl; alternatively, R^{15} and R^{16} can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring may be saturated, unsaturated or aromatic;

each of R^{17} and R^{53} is C_{1-5} alkyl, phenyl, or C_{1-5} heterocyclyl;

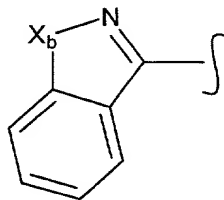
each of R^{54} and R^{55} is independently hydrogen, C_{1-5} alkyl, C_{2-5} alkenyl, phenyl, benzyl, or C_{1-5} heterocyclyl;

alternatively, R^{54} and R^{55} can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring may be saturated, unsaturated or aromatic;

W represents SO_2 , $C=O$, CHR^{20} , or a covalent bond; or W and R^1 , taken together with the 6-membered ring to which they are both attached, form one of the following two formulae:



(I)(a)



(I)(b)

wherein X_a is O, S, or N; and X_b is O, S or SO_2 ;

R^{20} is hydrogen, C_{1-5} alkyl, phenyl, benzyl, naphthyl, or C_{1-5} heterocyclyl;

R^{42} is hydrogen, C_{1-5} alkyl, C_{3-5} alkenyl, phenyl, naphthyl, C_{1-5} heterocyclyl, C_{2-8} acyl, aroyl, $R^{45}OC=O$, $R^{46}R^{47}NC=O$, $R^{45}SO$, $R^{45}SO_2$, or $R^{46}R^{47}NSO_2$;

R^{43} is hydrogen, C_{1-5} alkyl, C_{3-5} alkenyl, phenyl, or C_{1-5} heterocyclyl;
alternatively, R^{42} and R^{43} can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring may be saturated, unsaturated or aromatic;

R^{44} is C_{1-5} alkyl, C_{2-5} alkenyl, phenyl, naphthyl, or C_{1-5} heterocyclyl;

R^{48} is hydrogen, C_{1-5} alkyl, C_{3-5} alkenyl, phenyl, naphthyl, C_{1-5} heterocyclyl, C_{2-8} acyl, aroyl, $R^{50}OC=O$, $R^{51}R^{52}NC=O$, $R^{50}SO$, $R^{50}SO_2$, or $R^{51}R^{52}NSO_2$;

R^{49} is hydrogen, C_{1-5} alkyl, C_{3-5} alkenyl, phenyl, or C_{1-5} heterocyclyl;
alternatively, R^{48} and R^{49} can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring may be saturated, unsaturated or aromatic; and

wherein each of the above hydrocarbonyl or heterocarbonyl groups, unless otherwise indicated, and in addition to any specified substituents, is optionally and independently substituted with between 1 and 3 substituents selected from methyl, halomethyl, hydroxymethyl, halo, hydroxy, amino, nitro, cyano, C_{1-5} alkyl, C_{1-5} alkoxy, $-COOH$, C_{2-6} acyl, $[di(C_{1-4} \text{ alkyl})amino]C_{2-5}$ alkylene, $[di(C_{1-4} \text{ alkyl})amino] C_{2-5}$ alkyl-NH-CO-, and C_{1-5} haloalkoxy;

or a pharmaceutically acceptable salt, ester, or amide thereof, including a stereoisomeric form thereof.

The disclosed compounds are high-affinity inhibitors of the proteolytic activity of human cathepsin S. For use in medicine, the preparation of pharmaceutically acceptable salts of compounds of formula (I) may be desirable.

Certain compounds of the present invention may have one stereogenic atom and may exist as two enantiomers. Certain compounds of the present invention may have two or more stereogenic atoms and may further exist as

diastereomers. It is to be understood by those skilled in the art that all such stereoisomers and mixtures thereof in any proportion are encompassed within the scope of the present invention.

5 Another aspect of the invention provides pharmaceutical compositions comprising a compound of formula (I) and a pharmaceutically acceptable carrier. A further embodiment of the invention is a process for making a pharmaceutical composition comprising mixing a disclosed compound as described above, with a suitable pharmaceutically acceptable carrier.

10 The invention also contemplates pharmaceutical compositions comprising more than one compound of formula (I) and compositions comprising a compound of formula (I) and another pharmaceutically active agent.

15 The invention features a method of treating disorders or conditions mediated by the cathepsin S enzyme, in a subject in need thereof, comprising administering to the subject a therapeutically effective amount of any of the compounds or pharmaceutical compositions described above. If more than
20 one active agent is administered, the therapeutically effective amount may be a jointly effective amount. The compounds described herein inhibit the protease activity of human cathepsin S, an enzyme involved in the immune response. In preferred embodiments, cathepsin S inhibition is selective. As such, the disclosed compounds and compositions are useful in the prevention, inhibition,
25 or treatment of autoimmune diseases such as lupus, rheumatoid arthritis, and asthma, and for the prevention, inhibition, or treatment of tissue transplant rejection.

30 Additional features and advantages of the invention will become apparent from the detailed description below, including examples, and the appended claims.

Detailed Description of the Invention

The invention features pyrazole compounds of formula (I), methods of
5 making them, compositions containing them, and methods of using them to
treat diseases and conditions, including those mediated by Cathepsin S.

A. Terms

10 The following terms are defined below and by their usage throughout
this disclosure.

“Alkyl” includes optionally substituted straight chain and branched
hydrocarbons with at least one hydrogen removed to form a radical group.
15 Alkyl groups include methyl, ethyl, propyl, isopropyl, butyl, isobutyl, t-butyl, 1-
methylpropyl, pentyl, isopentyl, sec-pentyl, hexyl, heptyl, octyl, and so on.
Alkyl includes cycloalkyl, such as cyclopropyl, cyclobutyl, cyclopentyl, and
cyclohexyl.

20 “Alkenyl” includes optionally substituted straight chain and branched
hydrocarbon radicals as above with at least one carbon-carbon double bond
(sp^2). Alkenyls include ethenyl (or vinyl), prop-1-enyl, prop-2-enyl (or allyl),
isopropenyl (or 1-methylvinyl), but-1-enyl, but-2-enyl, butadienyls, pentenyls,
hexa-2,4-dienyl, and so on. Hydrocarbon radicals having a mixture of double
25 bonds and triple bonds, such as 2-penten-4-ynyl, are grouped as alkynyls
herein. Alkenyl includes cycloalkenyl. *Cis* and *trans* or (E) and (Z) forms are
included within the invention.

30 “Alkynyl” includes optionally substituted straight chain and branched
hydrocarbon radicals as above with at least one carbon-carbon triple bond (sp).
Alkynyls include ethynyl, propynyls, butynyls, and pentynyls. Hydrocarbon
radicals having a mixture of double bonds and triple bonds, such as 2-penten-
4-ynyl, are grouped as alkynyls herein. Alkynyl does not include cycloalkynyl.

“Alkoxy” includes an optionally substituted straight chain or branched alkyl group with a terminal oxygen linking the alkyl group to the rest of the molecule. Alkoxy includes methoxy, ethoxy, propoxy, isopropoxy, butoxy, t-butoxy, pentoxy and so on. “Aminoalkyl”, “thioalkyl”, and “sulfonylalkyl” are analogous to alkoxy, replacing the terminal oxygen atom of alkoxy with, respectively, NH (or NR), S, and SO₂. Heteroalkyl includes alkoxy, aminoalkyl, thioalkyl, and so on.

“Aryl” includes phenyl, naphthyl, biphenyl, tetrahydronaphthyl, and so on, any of which may be optionally substituted. Aryl also includes arylalkyl groups such as benzyl, phenethyl, and phenylpropyl. Aryl includes a ring system containing an optionally substituted 6-membered carbocyclic aromatic ring, said system may be bicyclic, bridge, and/or fused. The system may include rings that are aromatic, or partially or completely saturated. Examples of ring systems include indenyl, pentalenyl, 1-4-dihydronaphthyl, indanyl, benzimidazolyl, benzothiophenyl, indolyl, benzofuranyl, isoquinoliny, and so on.

“Heterocyclyl” includes optionally substituted aromatic and nonaromatic rings having carbon atoms and at least one heteroatom (O, S, N) or heteroatom moiety (SO₂, CO, CONH, COO) in the ring. Unless otherwise indicated, a heterocyclic radical may have a valence connecting it to the rest of the molecule through a carbon atom, such as 3-furyl or 2-imidazolyl, or through a heteroatom, such as N-piperidyl or 1-pyrazolyl. Preferably a monocyclic heterocyclyl has between 4 and 7 ring atoms, or between 5 and 6 ring atoms; there may be between 1 and 5 heteroatoms or heteroatom moieties in the ring, and preferably between 1 and 3. A heterocyclyl may be saturated, unsaturated, aromatic (e.g., heteroaryl), nonaromatic, or fused.

Heterocyclyl also includes fused, e.g., bicyclic, rings, such as those optionally condensed with an optionally substituted carbocyclic or heterocyclic five- or six-membered aromatic ring. For example, “heteroaryl” includes an

optionally substituted six-membered heteroaromatic ring containing 1, 2 or 3 nitrogen atoms condensed with an optionally substituted five- or six-membered carbocyclic or heterocyclic aromatic ring. Said heterocyclic five- or six-membered aromatic ring condensed with the said five- or six-membered aromatic ring may contain 1, 2 or 3 nitrogen atoms where it is a six-membered ring, or 1, 2 or 3 heteroatoms selected from oxygen, nitrogen and sulfur where it is a five-membered ring.

Examples of heterocycllys include thiazoylyl, furyl, pyranlyl, isobenzofuranyl, pyrrolyl, imidazolyl, pyrazolyl, isothiazolyl, isoxazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, indolizinyll, isoindolyl, indolyl, indazolyl, purinyl, quinolyl, furazanyl, pyrrolidinyl, pyrrolinyl, imdazolidinyl, imidazolinyll, pyrazolidinyl, pyrazolinyl, piperidyl, piperazinyl, indolinyll, and morpholinyl. For example, preferred heterocycllys or heterocyclic radicals include morpholinyl, piperazinyl, pyrrolidinyl, pyridyl, cyclohexylimino, cycloheptylimino, and more preferably, piperidyl.

Examples illustrating heteroaryl are thienyl, furanyl, pyrrolyl, imidazolyl, oxazolyl, thiazolyl, benzothienyl, benzofuranyl, benzimidazolyl, benzoxazolyl, benzothiazolyl.

"Acyl" refers to a carbonyl moiety attached to either a hydrogen atom (i.e., a formyl group) or to an optionally substituted alkyl or alkenyl chain, or heterocyclyl.

"Halo" or "halogen" includes fluoro, chloro, bromo, and iodo, and preferably chloro or bromo as a substituent.

"Alkanediyl" or "alkylene" represents straight or branched chain optionally substituted bivalent alkane radicals such as, for example, methylene, ethylene, propylene, butylene, pentylene or hexylene.

"Alkenediyl" represents, analogous to the above, straight or branched chain optionally substituted bivalent alkene radicals such as, for example, propenylene, butenylene, pentenylene or hexenylene. In such radicals, the carbon atom linking a nitrogen preferably should not be unsaturated.

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"Aroyl" refers to a carbonyl moiety attached to an optionally substituted aryl or heteroaryl group, wherein aryl and heteroaryl have the definitions provided above. In particular, benzoyl is phenylcarbonyl.

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As defined herein, two radicals, together with the atom(s) to which they are attached may form an optionally substituted 4- to 7-, 5- to 7-, or a 5- to 6-membered ring carbocyclic or heterocyclic ring, which ring may be saturated, unsaturated or aromatic. Said rings may be as defined above in the Summary of the Invention section. Particular examples of such rings are as follows in the

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"Pharmaceutically acceptable salts, esters, and amides" include carboxylate salts (e.g., C₁₋₈ alkyl, cycloalkyl, aryl, heteroaryl, or non-aromatic heterocyclic) amino acid addition salts, esters, and amides which are within a reasonable benefit/risk ratio, pharmacologically effective and suitable for contact with the tissues of patients without undue toxicity, irritation, or allergic response. Representative salts include hydrobromide, hydrochloride, sulfate, bisulfate, nitrate, acetate, oxalate, valerate, oleate, palmitate, stearate, laurate, borate, benzoate, lactate, phosphate, tosylate, citrate, maleate, fumarate, succinate, tartrate, naphthylate, mesylate, glucoheptonate, lactobionate, and laurylsulfonate. These may include alkali metal and alkali earth cations such as sodium, potassium, calcium, and magnesium, as well as non-toxic ammonium, quaternary ammonium, and amine cations such as tetramethyl ammonium, methylamine, trimethylamine, and ethylamine. See example, S.M. Berge, et al., "Pharmaceutical Salts," J. Pharm. Sci., 1977, 66:1-19 which is incorporated herein by reference. Representative pharmaceutically acceptable amides of the invention include those derived from ammonia, primary C₁₋₈ alkyl amines and secondary di (C₁₋₈ alkyl) amines. Secondary amines include 5- or

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6-membered heterocyclic or heteroaromatic ring moieties containing at least one nitrogen atom and optionally between 1 and 2 additional heteroatoms. Preferred amides are derived from ammonia, C₁₋₃ alkyl primary amines, and di (C₁₋₂ alkyl)amines. Representative pharmaceutically acceptable esters of the invention include C₁₋₇ alkyl, C₅₋₇ cycloalkyl, phenyl, and phenyl(C₁₋₆)alkyl esters. Preferred esters include methyl esters.

"Patient" or "subject" includes mammals such as humans and animals (dogs, cats, horses, rats, rabbits, mice, non-human primates) in need of observation, experiment, treatment or prevention in connection with the relevant disease or condition. Preferably, the patient or subject is a human.

"Composition" includes a product comprising the specified ingredients in the specified amounts as well as any product which results directly or indirectly from combinations of the specified ingredients in the specified amounts.

"Therapeutically effective amount" or "effective amount" means that amount of active compound or pharmaceutical agent that elicits the biological or medicinal response in a tissue system, animal or human that is being sought by a researcher, veterinarian, medical doctor or other clinician, which includes alleviation of the symptoms of the disease or disorder being treated.

Concerning the various radicals in this disclosure and in the claims, three general remarks are made. The first remark concerns valency. As with all hydrocarbon radicals, whether saturated, unsaturated or aromatic, and whether or not cyclic, straight chain, or branched, and also similarly with all heterocyclic radicals, each radical includes substituted radicals of that type and monovalent, bivalent, and multivalent radicals as indicated by the context of the claims. The context will indicate that the substituent is an alkylene or hydrocarbon radical with at least two hydrogen atoms removed (bivalent) or more hydrogen atoms removed (multivalent). An example of a bivalent radical linking two parts of the molecule is G in formula (I) which links two rings.

Second, radicals or structure fragments as defined herein are understood to include substituted radicals or structure fragments.

Hydrocarbyls include monovalent radicals containing carbon and hydrogen such as alkyl, alkenyl, alkynyl, cycloalkyl, and cycloalkenyl (whether aromatic or unsaturated), as well as corresponding divalent radicals such as alkylene, alkenylene, phenylene, and so on. Heterocarbyls include monovalent and divalent radicals containing carbon, hydrogen, and at least one heteroatom. Examples of monovalent heterocarbyls include acyl, acyloxy, alkoxyacyl, heterocyclyl, heteroaryl, aroyl, benzoyl, dialkylamino, hydroxyalkyl, and so on.

Using "alkyl" as an example, "alkyl" should be understood to include substituted alkyl having one or more substitutions, such as between 1 and 5, 1 and 3, or 2 and 4 substituents. The substituents may be the same (dihydroxy, dimethyl), similar (chlorofluoro), or different (chlorobenzyl- or aminomethyl-substituted). Examples of substituted alkyl include haloalkyl (such as fluoromethyl, chloromethyl, difluoromethyl, perchloromethyl, 2-bromoethyl, perfluoromethyl, and 3-iodocyclopentyl), hydroxyalkyl (such as hydroxymethyl, hydroxyethyl, 2-hydroxypropyl, aminoalkyl (such as aminomethyl, 2-aminoethyl, 3-aminopropyl, and 2-aminopropyl), nitroalkyl, alkylalkyl, and so on. A di(C₁₋₆ alkyl)amino group includes independently selected alkyl groups, to form, for example, methylpropylamino and isopropylmethylamino, in addition dialkylamino groups having two of the same alkyl group such as dimethyl amino or diethylamino.

Third, only stable compounds are intended. For example, where there is an NR'R" group, and R can be an alkenyl group, the double bond is at least one carbon removed from the nitrogen to avoid enamine formation. Similarly, where a dashed line is an optional sp² bond, if it is absent, the appropriate hydrogen atom(s) is(are) included.

Preferred substitutions for Ar include methyl, methoxy, fluoromethyl, difluoromethyl, perfluoromethyl (trifluoromethyl), 1-fluoroethyl, 2-fluoroethyl, ethoxy, fluoro, chloro, and bromo, and particularly methyl, bromo, chloro, perfluoromethyl, perfluoromethoxy, methoxy, and fluoro. Preferred substitution patterns for Ar or Ar₁ are 4-substituted or 3,4-disubstituted phenyl.

Compounds of the invention are further described in the next section.

B. Compounds

The invention features compounds of formula (I) as described in the
 5 Summary section.

Preferred compounds include those wherein:

- (a) R¹ is hydrogen, halogen, C₁₋₅ alkoxy, hydroxy, C₁₋₅ alkyl, cyano, nitro, R⁷R⁸N, C₂₋₈ acyl, or R¹⁰R¹¹NSO₂;
- 10 (b) R¹ is halogen, cyano, nitro, R⁷R⁸N, or R¹⁰R¹¹NSO₂;
- (c) R² is hydrogen;
- (d) each of R³ and R⁴ is independently hydrogen or C₁₋₃ alkyl;
- (e) one of R³ and R⁴ is hydrogen;
- (f) each of R³ and R⁴ is hydrogen;
- 15 (g) one of R⁵ and R⁶ is hydrogen and the other is a 5-7 membered carbocyclyl or heterocyclyl, optionally substituted;
- (h) R⁵ and R⁶ taken together form a six-membered heterocyclyl;
- (i) R⁵ and R⁶ taken together form pyridinyl, pyrimidinyl, or piperazinyl, optionally N-substituted with R⁴⁰O(C=O)(C=O)-, R⁴⁰SO₂, R⁴⁰NHCO₂, R⁴⁰(C=O)- or R⁴⁰N(C=O)-;
- 20 (j) each of R⁷, R⁸, R²¹, R²², R²⁴, R²⁵ is independently hydrogen or C₁₋₅ alkyl; or, independently, each of R⁷ and R⁸, R²¹ and R²², and R²⁴ and R²⁵ can be taken together to form an optionally substituted 4- to 7- membered heterocyclic ring, which ring
- 25 may be saturated, unsaturated or aromatic;
- (k) at least one of R⁷ and R⁸, R²¹ and R²², and R²⁴ and R²⁵, taken together, is morpholinyl, piperidinyl, or pyrrolidinyl;
- (l) R⁹, R²³, R²⁶, and R²⁷ is each independently hydrogen or C₁₋₅ alkyl;
- 30 (m) G is C₃₋₄ alkanediyl, optionally substituted with hydroxy, (L)-C₁₋₅ alkyloxy-, or [(L)-C₁₋₅ alkylene]amino-;
- (n) G is C₃ alkanediyl, optionally substituted with hydroxy, (L)-C₁₋₅ alkyloxy-, or [(L)-C₁₋₅ alkylene]amino-;

- (o) X is nitrogen;
- (p) Y is CR¹³;
- (q) Z is CR¹⁴;
- (r) X is CH;
- 5 (s) R¹² is hydrogen, R²²O(C=O)NH-, R²²NH(C=O)NH-, R²²SO₂NH-, R²³SO₂ or R²³SO₂-, and R¹³ is hydrogen, R⁴³O(C=O)NH-, R⁴³NH(C=O)NH-, R⁴³SO₂NH-, R⁴⁴SO₂ or R⁴⁴SO₂;
- (t) R¹⁴ is hydrogen, halogen, C₁₋₅ alkoxy, C₁₋₅ alkyl, cyano, nitro, R²⁵O(C=O)NH-, R²⁵NH(C=O)NH-, R²⁵SO₂NH or R²⁴R²⁵N;
- 10 (u) R¹⁴ is halogen, R²⁵O(C=O)NH-, R²⁵NH(C=O)NH-, R²⁵SO₂NH or R²⁴R²⁵N; Ar represents a monocyclic ring, optionally substituted with between 1 and 2 substituents selected independently from halogen, C₁₋₅ alkyl, cyano, nitro, R¹⁵R¹⁶N, CF₃ and OCF₃;
- 15 (v) Ar is a six membered ring substituted with between 1 and 2 substituents selected from halo, CF₃, OCF₃, said substituent or substituents being at the 4-position or at the 3- and 4-positions, respectively;
- (w) W is SO₂, C=O, or CHR²⁰;
- 20 (x) W is a covalent bond;
- (y) W and R¹ taken together are formula (I)(a);
- (z) W and R¹ taken together are formula (I)(b);
- (aa) one of R³ and R⁴ is hydrogen; Ar represents a monocyclic ring, optionally substituted with between 1 and 2 substituents selected from halogen, C₁₋₅ alkyl, cyano, nitro, R¹⁵R¹⁶N, CF₃ and OCF₃; R¹² is hydrogen, R²³SO₂ or R²³SO₂;-; R¹³ is hydrogen, R⁴⁴SO₂ or R⁴⁴SO₂;-; R¹⁴ is hydrogen, halogen, C₁₋₅ alkoxy, C₁₋₅ alkyl, cyano, nitro, or R²⁴R²⁵N; and G is C₃₋₄ alkanediyl, optionally substituted with hydroxy, C₁₋₃ alkyl, (L)-C₁₋₅ alkyloxy, or [(L)-C₁₋₅ alkylene]amino-;
- 25
- 30 (bb) each of R³ and R⁴ is hydrogen; Ar represents a six membered ring, optionally substituted with between 1 and 2 substituents selected from halogen, C₁₋₅ alkyl, cyano, nitro,

$R^{15}R^{16}N$, CF_3 and OCF_3 ; R^{12} is hydrogen, $R^{23}SO$, or $R^{23}SO_2$; R^{13} is hydrogen, $R^{44}SO$, or $R^{44}SO_2$; R^{14} is hydrogen, halogen, C_{1-5} alkoxy, C_{1-5} alkyl, cyano, nitro, or $R^{24}R^{25}N$; and G is C_3 alkanediyl, optionally substituted with hydroxy, (L)- C_{1-5} alkyloxy-, or (L)- C_{1-5} alkylamino;

(cc) Ar is phenyl; and

(dd) combinations of the above.

Specific preferred compounds include the examples herein, such as:

- 10 1-[4-(2-Amino-6-chloro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol ; 1-[3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-3-methyl-urea ; 1-[3-Chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-3-methyl-urea ;
- 15 3-Amino-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzoic acid methyl ester ; 3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenylamine ; 1-[2-(4-{3-[3-(4-Bromo-phenyl)-5-methanesulfonyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-3-chloro-phenyl]-3-methyl-urea ; 1-{3-[4-(2-Chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide ;
- 20 [3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-carbamic acid methyl ester ; 1-[3-(4-Benzo[d]isothiazol-3-yl)-piperazin-1-yl]-propyl]-3-(4-bromo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide ; 2-(4-{3-[5-Acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-3-nitro-benzoic acid methyl ester ;
- 25 1-[4-(2-Chloro-6-nitro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol ; 2-(4-{2-Hydroxy-3-[3-(4-iodo-phenyl)-5-
- 30

methanesulfonyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl)-piperazin-1-yl)-benzonitrile ; 3-(4-Bromo-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide ; 2-(4-{3-[5-Acetyl-3-(4-iodo-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile ; 2-(4-{3-[3-(4-Chloro-3-methyl-phenyl)-5-methanesulfonyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile ; 1-(3-(4-Chloro-3-methyl-phenyl)-1-{3-[4-(2,4-dimethyl-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone ; 1-{3-[4-(3,5-Dichloro-pyridin-4-yl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine ; 2-(4-{3-[5-Methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzonitrile; N-[3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-methanesulfonamide ; 3-(3,4-Dichloro-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide ; and 3-(4-Chloro-3-methyl-phenyl)-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic.

Furthermore, preferred compounds include those wherein Ar is selected from 4-trifluoromethylphenyl, 4-bromophenyl, 4-chlorophenyl, 4-chloro-3-methylphenyl and 3,4-dichlorophenyl.

More preferred compounds include the compounds in Examples 19, 27, and 33.

Related Compounds

The invention provides the disclosed compounds and closely related, pharmaceutically acceptable forms of the disclosed compounds, such as salts, esters, amides, acids, hydrates or solvated forms thereof; masked or protected forms; and racemic mixtures, or enantiomerically or optically pure forms. Related compounds also include compounds of the invention that have been modified to be detectable, e.g., isotopically labelled with ^{18}F for use as a probe in positron emission tomography (PET) or single-photon emission computed tomography (SPECT).

The invention also includes disclosed compounds having one or more functional groups (e.g., hydroxyl, amino, or carboxyl) masked by a protecting group. See, e.g., Greene and Wuts, Protective Groups in Organic Synthesis, 3rd ed., (1999) John Wiley & Sons, NY. Some of these masked or protected compounds are pharmaceutically acceptable; others will be useful as intermediates. Synthetic intermediates and processes disclosed herein, and minor modifications thereof, are also within the scope of the invention.

HYDROXYL PROTECTING GROUPS

Protection for the hydroxyl group includes methyl ethers, substituted
 methyl ethers, substituted ethyl ethers, substituted benzyl ethers, and silyl
 5 ethers.

Substituted Methyl Ethers

Examples of substituted methyl ethers include methoxymethyl,
 methylthiomethyl, *t*-butylthiomethyl, (phenyldimethylsilyl)methoxymethyl,
 10 benzyloxymethyl, *p*-methoxybenzyloxymethyl, (4-methoxyphenoxy)methyl,
 guaiacolmethyl, *t*-butoxymethyl, 4-pentenylloxymethyl, siloxymethyl, 2-
 methoxyethoxymethyl, 2,2,2-trichloroethoxymethyl, bis(2-chloroethoxy)methyl,
 2-(trimethylsilyl)ethoxymethyl, tetrahydropyranyl, 3-bromotetrahydropyranyl,
 tetrahydrothiopyranyl, 1-methoxycyclohexyl, 4-methoxytetrahydropyranyl, 4-
 15 methoxytetrahydrothiopyranyl, 4-methoxytetrahydrothiopyranyl S,S-dioxido, 1-
 [(2-chloro-4-methyl)phenyl]-4-methoxypiperidin-4-yl, 1,4-dioxan-2-yl,
 tetrahydrofuranyl, tetrahydrothiofuranyl and 2,3,3a,4,5,6,7,7a-octahydro-7,8,8-
 trimethyl-4,7-methanobenzofuran-2-yl.

20 Substituted Ethyl Ethers

Examples of substituted ethyl ethers include 1-ethoxyethyl, 1-(2-
 chloroethoxy)ethyl, 1-methyl-1-methoxyethyl, 1-methyl-1-benzyloxyethyl, 1-
 methyl-1-benzyloxy-2-fluoroethyl, 2,2,2-trichloroethyl, 2-trimethylsilylethyl, 2-
 (phenylselenyl)ethyl, *t*-butyl, allyl, *p*-chlorophenyl, *p*-methoxyphenyl, 2,4-
 25 dinitrophenyl, and benzyl.

Substituted Benzyl Ethers

Examples of substituted benzyl ethers include *p*-methoxybenzyl, 3,4-
 dimethoxybenzyl, *o*-nitrobenzyl, *p*-nitrobenzyl, *p*-halobenzyl, 2,6-
 30 dichlorobenzyl, *p*-cyanobenzyl, *p*-phenylbenzyl, 2- and 4-picolyl, 3-methyl-2-
 picolyl N-oxido, diphenylmethyl, *p*, *p'*-dinitrobenzhydryl, 5-dibenzosuberyl,

triphenylmethyl, α -naphthylmethyl, *p*-methoxyphenyldiphenylmethyl, di(*p*-methoxyphenyl)methyl, tri(*p*-methoxyphenyl)methyl, 4-(4'-bromophenoxy)phenyldiphenylmethyl, 4,4',4''-tris(4,5-dichlorophthalimidophenyl)methyl, 4,4',4''-tris(levulinoyloxyphenyl)methyl, 4,4',4''-tris(benzoyloxyphenyl)methyl, 3-(1-midazol-1-ylmethyl)bis(4',4''-dimethoxyphenyl)methyl, 1,1-bis(4-methoxyphenyl)-1'-pyrenylmethyl, 9-anthryl, 9-(9-phenyl)xanthenyl, 9-(9-phenyl-10-oxo)anthryl, 1,3-benzodithiolan-2-yl, and benzisothiazolyl S,S-dioxido.

Silyl Ethers

Examples of silyl ethers include trimethylsilyl, triethylsilyl, triisopropylsilyl, dimethylisopropylsilyl, diethylisopropylsilyl, dimethylthexylsilyl, *t*-butyldimethylsilyl, *t*-butyldiphenylsilyl, tribenzylsilyl, tri-*p*-xylylsilyl, triphenylsilyl, diphenylmethylsilyl, and *t*-butylmethoxyphenylsilyl.

Esters

In addition to ethers, a hydroxyl group may be protected as an ester. Examples of esters include formate, benzoylformate, acetate, chloroacetate, dichloroacetate, trichloroacetate, trifluoroacetate, methoxyacetate, triphenylmethoxyacetate, phenoxyacetate, *p*-chlorophenoxyacetate, *p*-phenylacetate, 3-phenylpropionate, 4-oxopentanoate(levulinate), 4,4-(ethylenedithio)pentanoate, pivaloate, adamantate, crotonate, 4-methoxycrotonate, benzoate, *p*-phenylbenzoate, 2,4,6-trimethylbenzoate(mesitoate)

Carbonates

Examples of carbonate protecting groups include methyl, 9-fluorenylmethyl, ethyl, 2,2,2-trichloroethyl, 2-(trimethylsilyl)ethyl, 2-(phenylsulfonyl)ethyl, 2-(triphenylphosphonio)ethyl, isobutyl, vinyl, allyl, *p*-nitrophenyl, benzyl, *p*-methoxybenzyl, 3,4-dimethoxybenzyl, *o*-nitrobenzyl, *p*-nitrobenzyl, S-benzyl thiocarbonate, 4-ethoxy-1-naphthyl, and methyl dithiocarbonate.

Assisted Cleavage

- Examples of assisted cleavage include 2-iodobenzoate, 4-azidobutyrate, 4-nitro-4-methylpentanoate, *o*-(dibromomethyl)benzoate, 2-formylbenzenesulfonate, 2-(methylthiomethoxy)ethyl carbonate, 4-(methylthiomethoxy)butyrate, and 2-(methylthiomethoxymethyl)benzoate.

Miscellaneous Esters

- Examples of miscellaneous esters include 2,6-dichloro-4-methylphenoxyacetate, 2,6-dichloro-4-(1,1,3,3-tetramethylbutyl)phenoxyacetate, 2,4-bis(1,1-dimethylpropyl)phenoxyacetate, chlorodiphenylacetate, isobutyrate, monosuccinoate, (E)-2-methyl-2-butenolate(tigloate), *o*-(methoxycarbonyl)benzoate, *p*-P-benzoate, α -naphthoate, nitrate, alkyl N,N,N',N'-tetramethylphosphorodiamidate, N-phenylcarbamate, borate, dimethylphosphinothioyl, and 2,4-dinitrophenylsulfenate.

Sulfonates

- Examples of sulfonates include sulfate, methanesulfonate(mesylate), benzyisulfonate, and tosylate.

25 AMINO PROTECTING GROUPS

Protection for the amino group includes carbamates, amides, and special –NH protective groups.

- Examples of carbamates include methyl and ethyl carbamates, substituted ethyl carbamates, assisted cleavage carbamates, photolytic cleavage carbamates, urea-type derivatives, and miscellaneous carbamates.

Carbamates

Examples of methyl and ethyl carbamates include methyl and ethyl, 9-fluorenylmethyl, 9-(2-sulfo)fluorenylmethyl, 9-(2,7-dibromo)fluorenylmethyl, 2,7-di-*t*-butyl-[9-(10,10-dioxo-10,10,10,10-tetrahydrothioxanthyl)]methyl, and 4-methoxyphenacyl.

Substituted Ethyl

Examples of substituted ethyl carbamates include 2,2,2-trichloroethyl, 2-trimethylsilylethyl, 2-phenylethyl, 1-(1-adamantyl)-1-methylethyl, 1,1-dimethyl-2-haloethyl, 1,1-dimethyl-2,2-dibromoethyl, 1,1-dimethyl-2,2,2-trichloroethyl, 1-methyl-1-(4-biphenyl)ethyl, 1-(3,5-di-*t*-butylphenyl)-1-methylethyl, 2-(2'- and 4'-pyridyl)ethyl, 2-(*N,N*-dicyclohexylcarboxamido)ethyl, *t*-butyl, 1-adamantyl, vinyl, allyl, 1-isopropylallyl, cinnamyl, 4-nitrocinnamyl, 8-quinolyl, *N*-hydroxypiperidiny, alkylidithio, benzyl, *p*-methoxybenzyl, *p*-nitrobenzyl, *p*-bromobenzyl, *p*-chlorobenzyl, 2,4-dichlorobenzyl, 4-methylsulfinylbenzyl, 9-anthrylmethyl and diphenylmethyl.

Assisted Cleavage

Examples of assisted cleavage include 2-methylthioethyl, 2-methylsulfonylethyl, 2-(*p*-toluenesulfonyl)ethyl, [2-(1,3-dithianyl)]methyl, 4-methylthiophenyl, 2,4-dimethylthiophenyl, 2-phosphonioethyl, 2-triphenylphosphonioisopropyl, 1,1-dimethyl-2-cyanoethyl, *m*-chloro-*p*-acyloxybenzyl, *p*-(dihydroxyboryl)benzyl, 5-benzisoxazolylmethyl, and 2-(trifluoromethyl)-6-chromonylmethyl.

Photolytic Cleavage

Examples of photolytic cleavage include *m*-nitrophenyl, 3,5-dimethoxybenzyl, *o*-nitrobenzyl, 3,4-dimethoxy-6-nitrobenzyl, and phenyl(*o*-nitrophenyl)methyl.

Urea-Type Derivatives

Examples of urea-type derivatives include phenothiazinyl-(10)-carbonyl derivative, N' -*p*-toluenesulfonylamino carbonyl, and N'-
 5 phenylaminothiocarbonyl.

Miscellaneous Carbamates

Examples of miscellaneous carbamates include *t*-amyl, *S*-benzyl
 10 thiocarbamate, *p*-cyanobenzyl, cyclobutyl, cyclohexyl, cyclopentyl, cyclopropylmethyl, *p*-decyloxybenzyl, diisopropylmethyl, 2,2-dimethoxycarbonylvinyl, *o*-(*N,N*-dimethylcarboxamido)benzyl, 1,1-dimethyl-3-(*N,N*-dimethylcarboxamido)propyl, 1,1-dimethylpropynyl, di(2-pyridyl)methyl, 2-furanylmethyl, 2-iodoethyl, isobornyl, isobutyl, isonicotinyl, *p*-(*p*'-
 15 methoxyphenylazo)benzyl, 1-methylcyclobutyl, 1-methylcyclohexyl, 1-methyl-1-cyclopropylmethyl, 1-methyl-1-(3,5-dimethoxyphenyl)ethyl, 1-methyl-1-(*p*-phenylazophenyl)ethyl, 1-methyl-1-phenylethyl, 1-methyl-1-(4-pyridyl)ethyl, phenyl, *p*-(phenylazo)benzyl, 2,4,6-tri-*t*-butylphenyl, 4-(trimethylammonium)benzyl, and 2,4,6-trimethylbenzyl.

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Examples of amides include:

Amides

N-formyl, N-acetyl, N-chloroacetyl, N-trichloroacetyl, N-trifluoroacetyl, N-
 25 phenylacetyl, N-3-phenylpropionyl, N-picolinoyl, N-3-pyridylcarboxamide, N-benzoylphenylalanyl derivative, N-benzoyl, N-*p*-phenylbenzoyl.

Assisted Cleavage

N-*o*-nitrophenylacetyl, N-*o*-nitrophenoxyacetyl, N-acetoacetyl, (N'-
 30 dithiobenzoyloxycarbonylamino)acetyl, N-3-(*p*-hydroxyphenyl)propionyl, N-3-(*o*-nitrophenyl)propionyl, N-2-methyl-2-(*o*-nitrophenoxy)propionyl, N-2-methyl-2-(*o*-phenylazophenoxy)propionyl, N-4-chlorobutyryl, N-3-methyl-3-nitrobutyryl, N-*o*-nitrocinnamoyl, N-acetylmethionine derivative, N-*o*-nitrobenzoyl, N-*o*-(benzoyloxymethyl)benzoyl, and 4,5-diphenyl-3-oxazolin-2-one.

Cyclic Imide Derivatives

5 N-phthalimide, N-dithiasuccinoyl, N-2,3-diphenylmaleoyl, N-2,5-dimethylpyrrolyl, N-1,1,4,4-tetramethyldisilylazacyclopentane adduct, 5-substituted 1,3-dimethyl-1,3,5-triazacyclohexan-2-one, 5-substituted 1,3-dibenzyl-1,3,5-triazacyclohexan-2-one, and 1-substituted 3,5-dinitro-4-pyridonyl.

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SPECIAL – NH PROTECTIVE GROUPS

Examples of special NH protective groups include

15 N-Alkyl and N-Aryl Amines

N-methyl, N-allyl, N-[2-(trimethylsilyl)ethoxy]methyl, N-3-acetoxypyrrol, N-(1-isopropyl-4-nitro-2-oxo-3-pyrrolin-3-yl), quaternary ammonium salts, N-benzyl, N-di(4-methoxyphenyl)methyl, N-5-dibenzosuberyl, N-triphenylmethyl, 20 N-(4-methoxyphenyl)diphenylmethyl, N-9-phenylfluorenyl, N-2,7-dichloro-9-fluorenylmethylene, N-ferrocenylmethyl, and N-2-picolylamine N'-oxide.

Imine Derivatives

25 N-1,1-dimethylthiomethylene, N-benzylidene, N-p-methoxybenzylidene, N-diphenylmethylene, N-[(2-pyridyl)mesityl]methylene, and N-(N',N'-dimethylaminomethylene).

30 PROTECTION FOR THE CARBONYL GROUP

Acyclic Acetals and Ketals

Examples of acyclic acetals and ketals include dimethyl, bis(2,2,2-trichloroethyl), dibenzyl, bis(2-nitrobenzyl) and diacetyl.

35

Cyclic Acetals and Ketals

- 5 Examples of cyclic acetals and ketals include 1,3-dioxanes, 5-methylene-1,3-dioxane, 5,5-dibromo-1,3-dioxane, 5-(2-pyridyl)-1,3-dioxane, 1,3-dioxolanes, 4-bromomethyl-1,3-dioxolane, 4-(3-butenyl)-1,3-dioxolane, 4-phenyl-1,3-dioxolane, 4-(2-nitrophenyl)-1,3-dioxolane, 4,5-dimethoxymethyl-1,3-dioxolane, O,O'-phenylenedioxy and 1,5-dihydro-3H-2,4-benzodioxepin.

10 Acyclic Dithio Acetals and Ketals

 Examples of acyclic dithio acetals and ketals include S,S'-dimethyl, S,S'-diethyl, S,S'-dipropyl, S,S'-dibutyl, S,S'-dipentyl, S,S'-diphenyl, S,S'-dibenzyl and S,S'-diacetyl.

15 Cyclic Dithio Acetals and Ketals

 Examples of cyclic dithio acetals and ketals include 1,3-dithiane, 1,3-dithiolane and 1,5-dihydro-3H-2,4-benzodithiepin.

20 Acyclic Monothio Acetals and Ketals

 Examples of acyclic monothio acetals and ketals include O-trimethylsilyl-S-alkyl, O-methyl-S-alkyl or -S-phenyl and O-methyl-S-2-(methylthio)ethyl.

25 Cyclic Monothio Acetals and Ketals

 Examples of cyclic monothio acetals and ketals include 1,3-oxathiolanes.

30

MISCELLANEOUS DERIVATIVES

O-Substituted Cyanohydrins

- 35 Examples of O-substituted cyanohydrins include O-acetyl, O-trimethylsilyl, O-1-ethoxyethyl and O-tetrahydropyranyl.

Substituted Hydrazones

Examples of substituted hydrazones include N,N-dimethyl and 2,4-dinitrophenyl.

5

Oxime Derivatives

Examples of oxime derivatives include O-methyl, O-benzyl and O-phenylthiomethyl.

10

Imines

Substituted Methylene Derivatives, Cyclic Derivatives

15

Examples of substituted methylene and cyclic derivatives include oxazolidines, 1-methyl-2-(1'-hydroxyalkyl)imidazoles, N,N'-dimethylimidazolidines, 2,3-dihydro-1,3-benzothiazoles, diethylamine adducts, and methylaluminum bis(2,6-di-*t*-butyl-4-methylphenoxide)(MAD)complex.

PROTECTION FOR THE CARBOXYL GROUP

Esters

5 Substituted Methyl Esters

Examples of substituted methyl esters include 9-fluorenylmethyl, methoxymethyl, methylthiomethyl, tetrahydropyranyl, tetrahydrofuranyl, methoxyethoxymethyl, 2-(trimethylsilyl)ethoxymethyl, benzyloxymethyl, 10 phenacyl, *p*-bromophenacyl, α -methylphenacyl, *p*-methoxyphenacyl, carboxamidomethyl, and N-phthalimidomethyl.

2-Substituted Ethyl Esters

15 Examples of 2-substituted ethyl esters include 2,2,2-trichloroethyl, 2-haloethyl, ω -chloroalkyl, 2-(trimethylsilyl)ethyl, 2-methylthioethyl, 1,3-dithianyl-2-methyl, 2-(*p*-nitrophenylsulfenyl)ethyl, 2-(*p*-toluenesulfonyl)ethyl, 2-(2'-pyridyl)ethyl, 2-(diphenylphosphino)ethyl, 1-methyl-1-phenylethyl, *t*-butyl, cyclopentyl, cyclohexyl, allyl, 3-buten-1-yl, 4-(trimethylsilyl)-2-buten-1-yl, 20 cinnamyl, α -methylcinnamyl, phenyl, *p*-(methylmercapto)phenyl and benzyl.

Substituted Benzyl Esters

Examples of substituted benzyl esters include triphenylmethyl, 25 diphenylmethyl, bis(*o*-nitrophenyl)methyl, 9-anthrylmethyl, 2-(9,10-dioxo)anthrylmethyl, 5-dibenzosuberyl, 1-pyrenylmethyl, 2-(trifluoromethyl)-6-chromylmethyl, 2,4,6-trimethylbenzyl, *p*-bromobenzyl, *o*-nitrobenzyl, *p*-nitrobenzyl, *p*-methoxybenzyl, 2,6-dimethoxybenzyl, 4-(methylsulfinyl)benzyl, 4-sulfobenzyl, piperonyl, 4-picolyl and *p*-P-benzyl.

30

Silyl Esters

Examples of silyl esters include trimethylsilyl, triethylsilyl, *t*-butyldimethylsilyl, *i*-propyldimethylsilyl, phenyldimethylsilyl and di-*t*- 35 butylmethylsilyl.

Activated Esters

Examples of activated esters include thiols.

5

Miscellaneous Derivatives

Examples of miscellaneous derivatives include oxazoles, 2-alkyl-1,3-oxazolines, 4-alkyl-5-oxo-1,3-oxazolidines, 5-alkyl-4-oxo-1,3-dioxolanes, ortho esters, phenyl group and pentaaminocobalt(III) complex.

10

Stannyl Esters

Examples of stannyl esters include triethylstannyl and tri-*n*-butylstannyl.

15

AMIDES AND HYDRAZIDES

Amides

Examples of amides include N,N-dimethyl, pyrrolidinyl, piperidinyl, 5,6-dihydrophenanthridinyl, o-nitroanilides, N-7-nitroindolyl, N-8-Nitro-1,2,3,4-tetrahydroquinolyl, and *p*-P-benzenesulfonamides.

20

Hydrazides

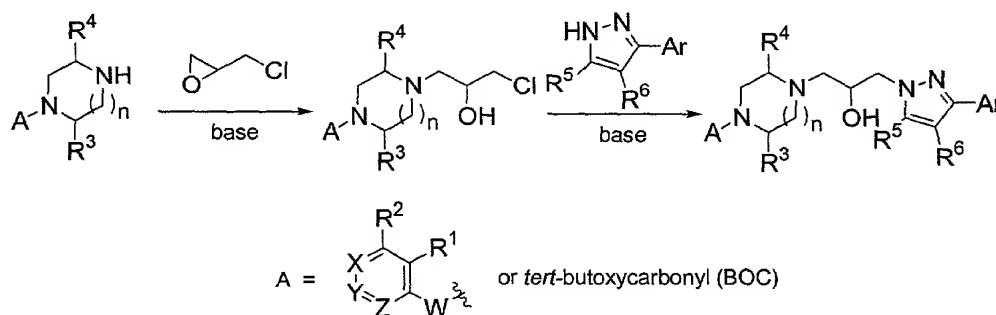
Examples of hydrazides include N-phenyl and N,N'-diisopropyl hydrazides.

25

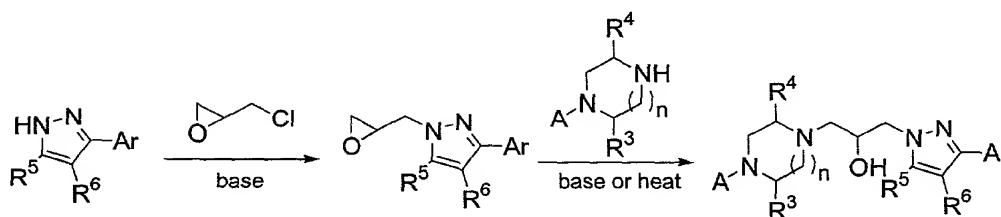
C. Synthesis

- The compounds of the present invention may be prepared by conventional synthetic organic chemistry and by matrix or combinatorial methods according to Schemes 1 to 11 below, and representative detailed Examples 1 to 24. Those of ordinary skill in the art will be able to modify and adapt the guidance provided herein to make the disclosed compounds.

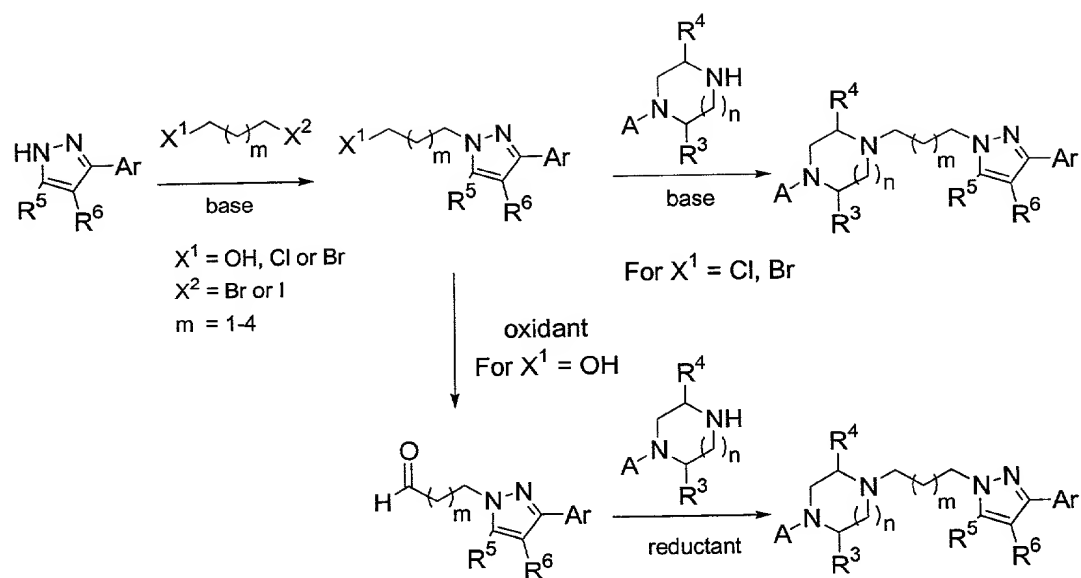
10 Scheme 1



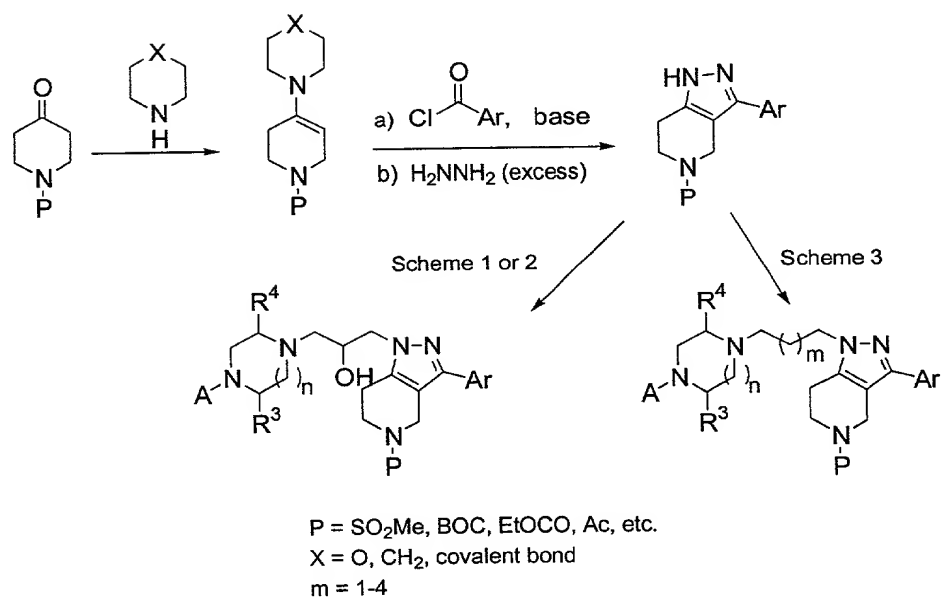
15 Scheme 2



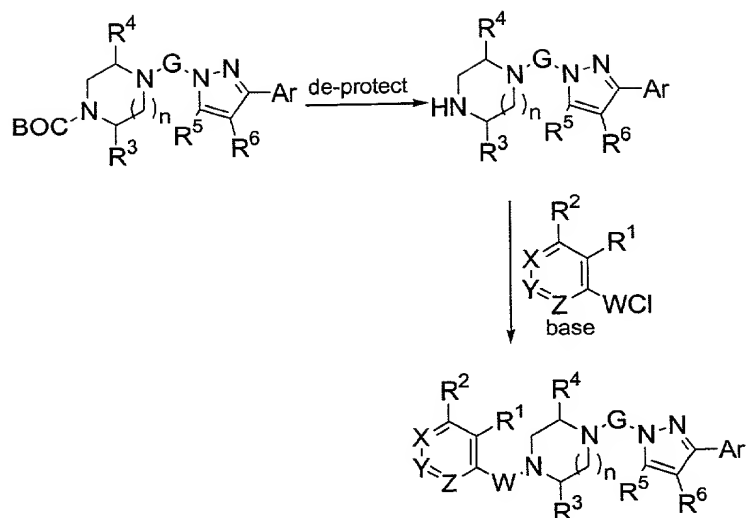
Scheme 3



5 Scheme 4

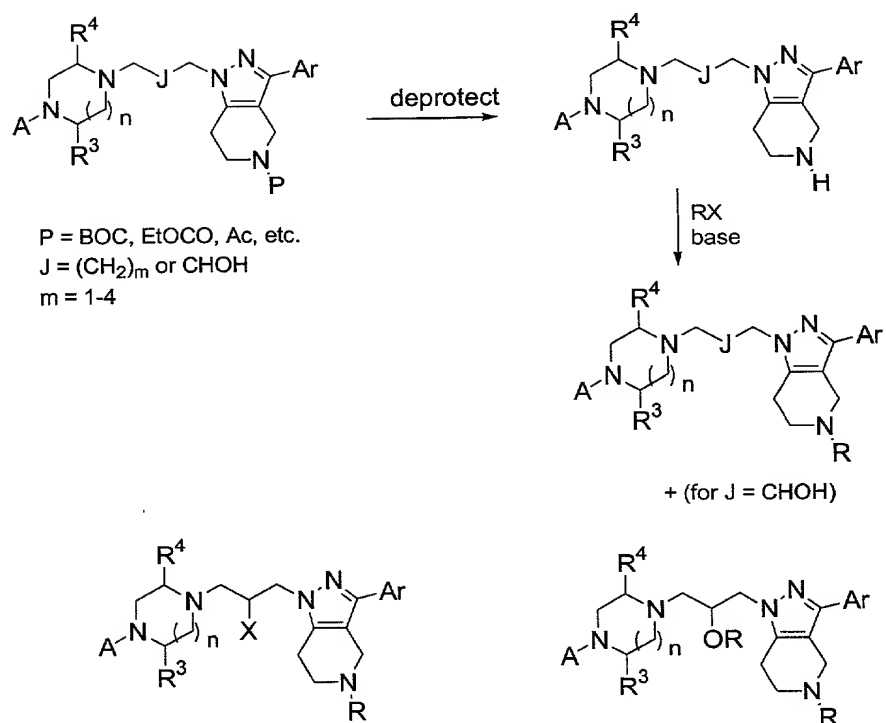


Scheme 5



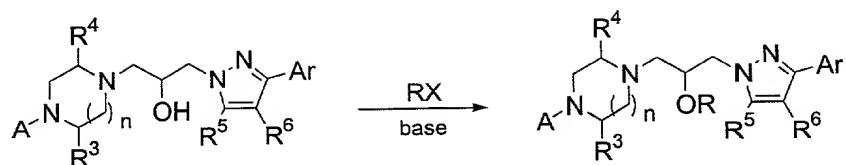
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Scheme 6



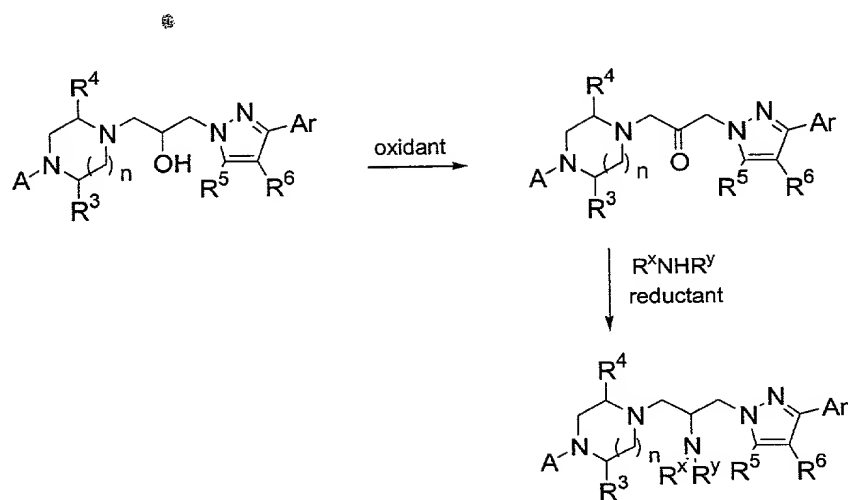
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Scheme 7



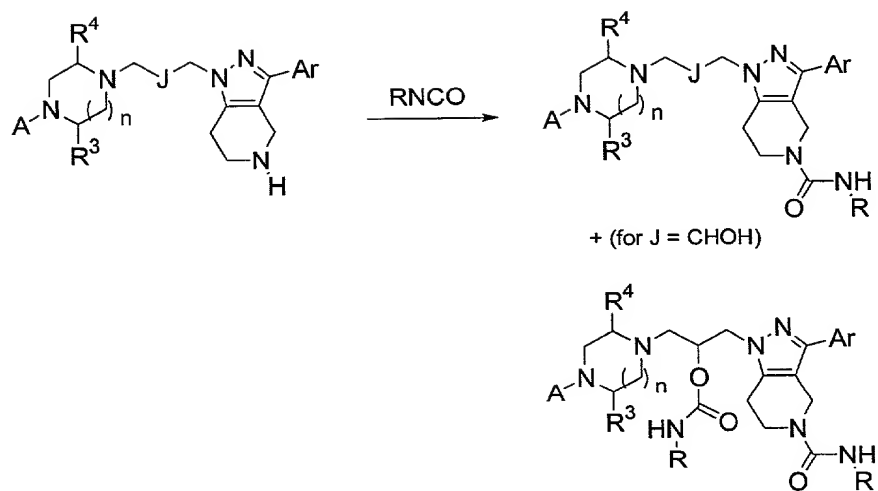
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Scheme 8

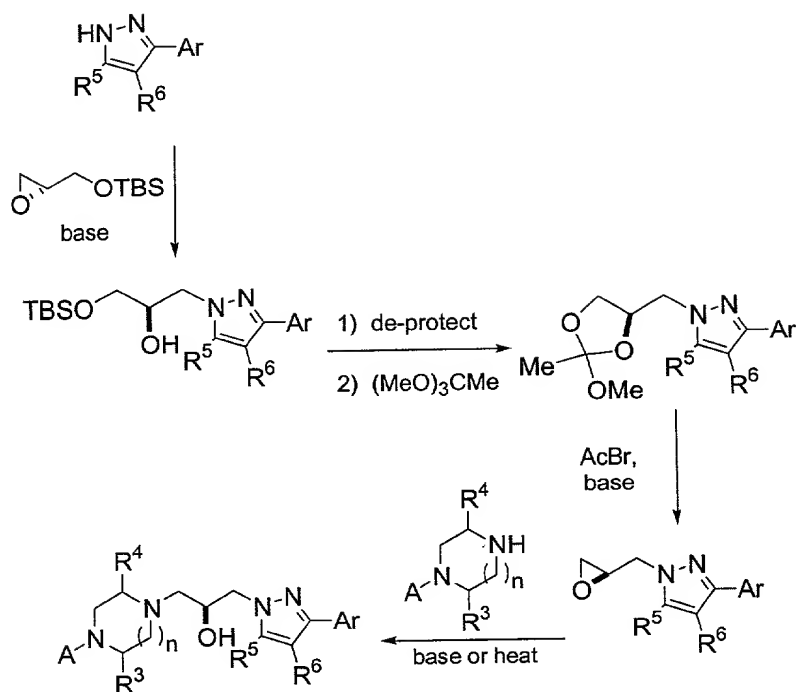


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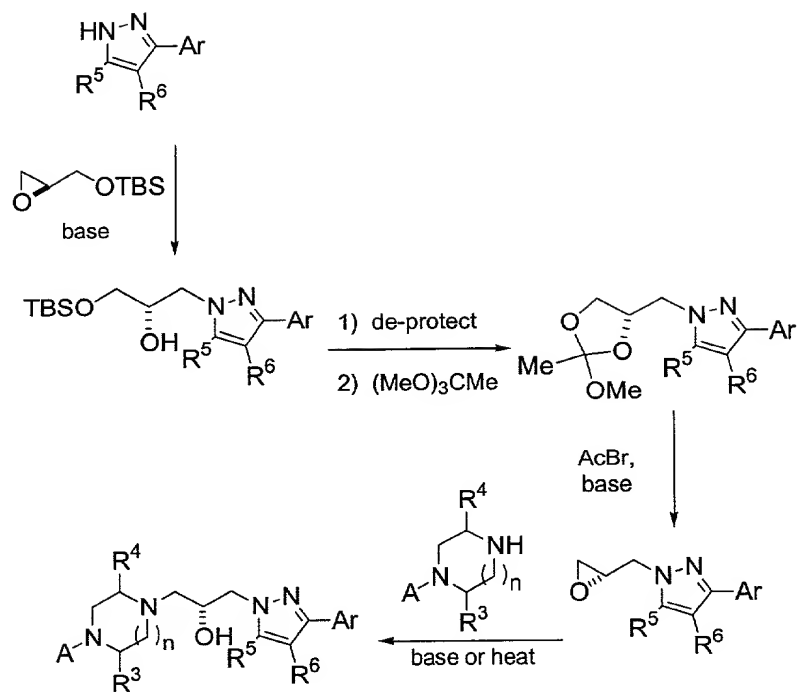
Scheme 9



Scheme 10



Scheme 11



D. Formulation and Administration

The present compounds inhibit the proteolytic activity of human cathepsin S and therefore are useful as a medicine especially in methods for treating patients suffering from disorders or conditions which are modulated or regulated by the inhibition of cathepsin S activity.

The invention features a method for treating a subject with a condition mediated by cathepsin S, said method comprising administering to the subject a therapeutically effective amount of a pharmaceutical composition comprising a compound of the invention. The invention also provides a method for inhibiting cathepsin S activity in a subject, wherein the method comprises administering to the subject a therapeutically effective amount of a pharmaceutical composition comprising a compound of the invention. A third method is a method for treating an autoimmune disease, or inhibiting the progression of an autoimmune disease, in a subject, said method comprising administering to the subject a therapeutically effective amount of a pharmaceutical composition comprising a disclosed compound. The autoimmune disease can be, for example, lupus, rheumatoid arthritis, or preferably, asthma. The invention also provides a method for treating or inhibiting the progression of tissue transplant rejection in a subject, the method comprising administering to the subject a therapeutically effective amount of a pharmaceutical composition comprising a compound of the invention. The administration step can occur before, during, and/or after a tissue transplant procedure.

In view of their inhibitory effect on the proteolytic activity of human cathepsin S the compounds of the present invention may be formulated into various pharmaceutical forms for administration purposes. To prepare these pharmaceutical compositions, an effective amount of a particular compound, in base or acid addition salt form, as the active ingredient is intimately mixed with a pharmaceutically acceptable carrier.

A carrier may take a wide variety of forms depending on the form of preparation desired for administration. These pharmaceutical compositions are desirably in unitary dosage form suitable, preferably, for oral administration or parenteral injection. For example, in preparing the compositions in oral dosage form, any of the usual pharmaceutical media may be employed. These include water, glycols, oils, alcohols and the like in the case of oral liquid preparations such as suspensions, syrups, elixirs and solutions; or solid carriers such as starches, sugars, kaolin, lubricants, binders, disintegrating agents and the like in the case of powders, pills, capsules and tablets. In view of their ease in administration, tablets and capsules represent the most advantageous oral dosage unit form, in which case solid pharmaceutical carriers are generally employed. For parenteral compositions, the carrier will usually comprise sterile water, at least in large part, though other ingredients, for example, to aid solubility, may be included. Injectable solutions, for example, may be prepared in which the carrier comprises saline solution, glucose solution or a mixture of saline and glucose solution. Injectable suspensions may also be prepared in which case appropriate liquid carriers, suspending agents and the like may be employed. In the compositions suitable for percutaneous administration, the carrier optionally comprises a penetration enhancing agent and/or a suitable wetting agent, optionally combined with suitable additives of any nature in minor proportions, which additives do not cause a significant deleterious effect to the skin. Such additives may facilitate the administration to the skin and/or may be helpful for preparing the desired compositions. These compositions may be administered in various ways, e.g., as a transdermal patch, as a spot-on, as an ointment. Acid addition salts of the compounds of formula (I), due to their increased water solubility over the corresponding base form, are more suitable in the preparation of aqueous compositions.

It is especially advantageous to formulate the aforementioned pharmaceutical compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used in the specification herein refers to physically discrete units suitable as unitary dosages, each unit containing a predetermined quantity of active ingredient calculated to produce

the desired therapeutic effect in association with the required pharmaceutical carrier. Examples of such dosage unit forms are tablets (including scored or coated tablets), capsules, pills, powder packets, wafers, injectable solutions or suspensions, teaspoonfuls, tablespoonfuls and the like, and segregated

5 multiples thereof.

Pharmaceutically acceptable acid addition salts include the therapeutically active non-toxic acid addition salt forms which the disclosed compounds are able to form. The latter can conveniently be obtained by treating the base

10 form with an appropriate acid. Appropriate acids comprise, for example, inorganic acids such as hydrohalic acids, e.g. hydrochloric or hydrobromic acid; sulfuric; nitric; phosphoric and the like acids; or organic acids such as, for example, acetic, propanoic, hydroxyacetic, lactic, pyruvic, oxalic, malonic, succinic, maleic, fumaric, malic, tartaric, citric, methanesulfonic,

15 ethanesulfonic, benzenesulfonic, *p*-toluenesulfonic, cyclamic, salicylic, *p*-aminosalicylic, palmoic and the like acids. The term addition salt also comprises the solvates which the disclosed compounds, as well as the salts thereof, are able to form. Such solvates are for example hydrates, alcoholates and the like. Conversely the salt form can be converted by treatment with alkali

20 into the free base form.

Stereoisomeric form defines all the possible isomeric forms which the compounds of formula (I) may possess. Unless otherwise mentioned or indicated, the chemical designation of compounds denotes the mixture of all

25 possible stereochemically isomeric forms, said mixtures containing all diastereomers and enantiomers of the basic molecular structure. More in particular, stereogenic centers may have the (R)- or (S)-configuration; substituents on bivalent cyclic saturated radicals may have either the cis- or trans-configuration. The invention encompasses stereochemically isomeric

30 forms including diastereoisomers, as well as mixtures thereof in any proportion of the disclosed compounds. The disclosed compounds may also exist in their tautomeric forms. Such forms although not explicitly indicated in the above

and following formulae are intended to be included within the scope of the present invention.

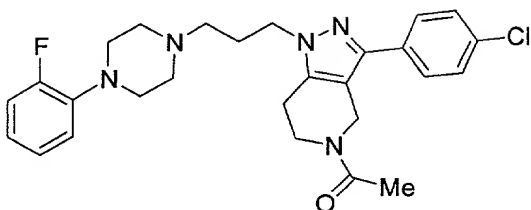
Those of skill in the treatment of disorders or conditions mediated by the cathepsin S enzyme could easily determine the effective daily amount from the test results presented hereinafter and other information. In general it is contemplated that a therapeutically effective dose would be from 0.001 mg/kg to 5 mg/kg body weight, more preferably from 0.01 mg/kg to 0.5 mg/kg body weight. It may be appropriate to administer the therapeutically effective dose as two, three, four or more sub-doses at appropriate intervals throughout the day. Said sub-doses may be formulated as unit dosage forms, for example, containing 0.05 mg to 250 mg, and in particular 0.5 to 50 mg of active ingredient per unit dosage form. Examples include 2 mg, 4 mg, 7 mg, 10 mg, 15 mg, 25 mg, and 35 mg dosage forms. Compounds of the invention may also be prepared in time-release or subcutaneous or transdermal patch formulations. Disclosed compound may also be formulated as a spray or other topical or inhalable formulations.

The exact dosage and frequency of administration depends on the particular compound of formula (I) used, the particular condition being treated, the severity of the condition being treated, the age, weight and general physical condition of the particular patient as well as other medication the patient may be taking, as is well known to those skilled in the art. Furthermore, it is evident that said effective daily amount may be lowered or increased depending on the response of the treated patient and/or depending on the evaluation of the physician prescribing the compounds of the instant invention. The effective daily amount ranges mentioned herein are therefore only guidelines.

The next section includes detailed information relating to the preparation, characterization, and use of the disclosed compounds.

E. Examples

5 EXAMPLE 1



1-(3-(4-Chloro-phenyl)-1-{3-[4-(2-fluoro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone.

10

A. 1-[3-(4-Chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

To a stirred solution of 50 g (0.35 mol) of *N*-acetyl-4-piperidone and 31 g (0.35 mol) of morpholine in benzene (350 mL) was added a catalytic amount (~ 0.25 g) of *p*-toluenesulfonic acid. The mixture was heated to reflux for 10 h with a Dean-Stark trap. The solvent was removed under reduced pressure to give a brown oil. The crude product was diluted with CH₂Cl₂ (175 mL) and 50.0 mL (0.35 mol) of Et₃N was added. The mixture was cooled to 0 °C and a solution of 45.0 mL (0.35 mol) of 4-chlorobenzoyl chloride in CH₂Cl₂ (50 mL) was added slowly by dropping funnel over 1 h. The mixture was allowed to warm to room temperature and stirred overnight. The reaction was then diluted with 1 *N* HCl (150 mL) and stirred vigorously for 3 h. The aqueous layer was extracted with CH₂Cl₂ (3 x 250 mL) and the combined extracts were dried over Na₂SO₄ and the solvent was removed under reduced pressure. The crude oil was diluted with EtOH (350 mL) and cooled to 0 °C. To this stirred solution was slowly added 33.0 mL (1.06 mol) of hydrazine and the mixture was allowed to warm to room temperature and stir overnight during which time a white precipitate formed. The volume of the reaction was reduced to ~150 mL and EtOAc (750 mL) was added to the mixture. The suspension was stirred vigorously for 2 h and was filtered then washed with EtOAc (2 x 200 mL) and dried under vacuum to afford 41.4 g (42% over 3 steps) of a pale yellow solid. TLC (silica,

5% MeOH/CH₂Cl₂): R_f = 0.3. MS (electrospray), *m/z* calculated for C₁₄H₁₄ClN₃O [M+H]⁺ 276.08, observed 276.0. ¹H NMR (400 MHz, CDCl₃, a mixture of amide rotamers): 7.65 (d, *J* = 8.4 Hz, 2H), 7.64 (d, *J* = 9.3 Hz, 2H), 7.58 (d, *J* = 10.5 Hz, 2H), 7.55 (d, *J* = 8.5 Hz, 2H), 4.94 (s, 2H), 4.78 (s, 2H), 4.08 (t, *J* = 5.9 Hz, 2H), 3.90 (t, *J* = 5.8 Hz, 2H), 3.02 (t, *J* = 5.8 Hz, 2H), 2.96 (t, *J* = 5.9 Hz, 2H), 2.36 (s, 3H), 2.31 (s, 3H).

B. 1-[3-(4-Chloro-phenyl)-1-(3-chloro-propyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

Cs₂CO₃ (2.66 g, 8.2 mmol) was added to a solution of 1-[3-(4-chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (1.0 g, 5.4 mmol) in DMF (10 mL) and stirred for 15 min. 1-Bromo-3-chloropropane (1.28 g, 8.2 mmol) was added and stirred under N₂ at room temperature for 36 h. Water (50 mL) was added to the reaction and stirred for 5 min. The product precipitated out. The aqueous portion was decanted and water was added to the residue and decanted again. The semisolid was taken up in CH₂Cl₂ and passed through a short plug of SiO₂ (5% MeOH/EtOAc) to obtain 1.06 g (83%) of a pale yellow semisolid. MS (electrospray): exact mass calculated for C₁₇H₁₉Cl₂N₃O, 351.09; *m/z* found, 352.0 [M+H]⁺. ¹H NMR (500 MHz, CDCl₃, a mixture of 1:1 rotamers): 7.60 (d, *J* = 8.3 Hz, 1H), 7.53 (d, *J* = 8.3 Hz, 1H), 7.40 (d, *J* = 8.3 Hz, 1H), 7.36 (d, *J* = 8.3 Hz, 1H), 4.77 (s, 1H), 4.61 (s, 1H), 4.20 (t, *J* = 6.2 Hz, 2H), 3.94 (t, *J* = 5.8 Hz, 1H), 3.76 (t, *J* = 5.8 Hz, 1H), 3.52 (q, *J* = 6.1 Hz, 2H), 2.84 (t, *J* = 5.5 Hz, 1H), 2.77 (t, *J* = 5.6 Hz, 1H), 2.37 (sextet, *J* = 6.1 Hz, 2H), 2.21 (s, 1.5 H), 2.16 (s, 1.5 H).

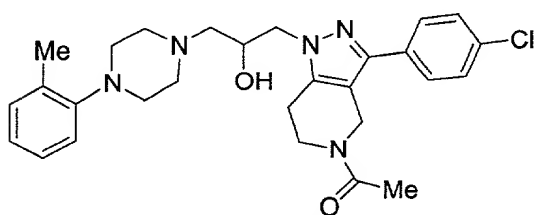
C. 1-(3-(4-Chloro-phenyl)-1-{3-[4-(2-fluoro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone.

1-[3-(4-Chloro-phenyl)-1-(3-chloro-propyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (0.053 g, 0.15 mmol) was dissolved in CH₃CN (0.5 mL) and a solution of 1-(2-fluorophenyl)piperazine (0.053 g, 0.30 mmol) in CH₃CN (0.5 mL) was added, followed by K₂CO₃ (0.031 g, 0.22 mmol) and Bu₄NI (0.018 g, 0.05 mmol). The mixture was stirred at room temperature for 7 d.

Preparative TLC (silica, 5% MeOH/EtOAc) afforded 30 mg (41%) of the title

compound. MS (electrospray): exact mass calculated for $C_{27}H_{31}ClFN_5O$, 495.22; m/z found, 496.3 $[M+H]^+$. 1H NMR (500 MHz, $CDCl_3$, a mixture of 1:1 rotamers): 7.60 (d, $J = 8.3$ Hz, 1H), 7.54 (d, $J = 8.3$ Hz, 1H), 7.39 (d, $J = 8.3$ Hz, 1H), 7.35 (d, $J = 8.3$ Hz, 1H), 7.06-6.90 (m, 4H), 4.77 (s, 1H), 4.60 (s, 1H), 4.10 (t, $J = 6.8$ Hz, 2H), 3.92 (t, $J = 5.7$ Hz, 1H), 3.74 (t, $J = 5.7$ Hz, 1H), 3.08 (br s, 4H), 2.83 (t, $J = 5.6$ Hz, 1H), 2.77 (t, $J = 5.7$ Hz, 1H), 2.58 (br s, 4H), 2.41-2.38 (m, 2H), 2.19 (s, 1.5H), 2.13 (s, 1.5H), 2.10-2.07 (m, 2H).

10 EXAMPLE 2



1-{3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone.

15

A. 1-[3-(4-Chloro-phenyl)-1-oxiranylmethyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

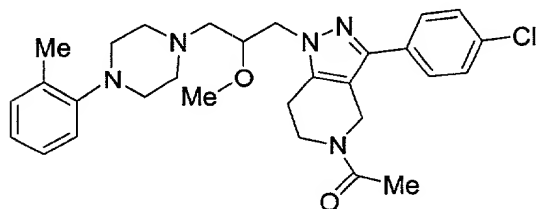
To a stirred solution of 1.00 g (3.63 mmol) of 1-[3-(4-chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone and 2.85 mL (36.3 mmol) of epichlorohydrin was added 1.30 g (3.99 mmol) of solid Cs_2CO_3 . The reaction was stirred for 48 h and the solvent was removed under reduced pressure. The residue was then diluted with H_2O (50 mL) and EtOAc (50 mL). The layers were separated, and the organic layer was washed with H_2O (25 mL) and brine (25 mL), dried over Na_2SO_4 and the solvent was removed under reduced pressure. Purification by flash chromatography (silica, 0-15% acetone/ CH_2Cl_2) afforded 0.72 g (60%) of a white solid. TLC (silica, 5% MeOH/ CH_2Cl_2): $R_f = 0.5$. MS (electrospray): m/z calculated for $C_{17}H_{18}ClN_3O_2$ $[M+H]^+$, 332.11, observed 332.0. 1H NMR (400 MHz, $CDCl_3$, a mixture of amide rotamers): 7.60 (d, $J = 8.6$ Hz, 2H), 7.54 (d, $J = 8.4$ Hz, 2H), 7.40 (d, $J = 8.6$ Hz, 2H), 7.36 (d, $J = 8.4$ Hz, 2H), 4.80 and 4.73 (A and B of AB quartet, $J_{ab} = 15.8$ Hz, 2H), 4.60 (s, 2H), 4.47 (dd, $J = 15.3, 2.5$ Hz, 1H), 4.42 (dd, $J = 15.0, 2.7$ Hz, 1H), 4.11 (dd, $J =$

5.3, 2.5 Hz, 1H), 4.08 (dd, $J = 5.1, 3.3$ Hz, 1H), 3.99-3.85 (m, 2H), 3.73 (dt, $J = 5.9, 1.8$ Hz, 2H), 3.37 (m, 2H), 2.87-2.80 (m, 3H), 2.80-2.69 (m, 3H), 2.53 (dd, $J = 4.7, 2.5$ Hz, 1H), 2.48 (dd, $J = 4.6, 2.6$, 1H), 2.19 (s, 3H), 2.15 (s, 3H).

5 B. 1-[3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl)-piperazin-1-yl]-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A solution of of 1-[3-(4-chloro-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (0.8 g, 2.42 mmol) in CH_2Cl_2 (12 mL) was treated with ytterbium(III) triflate (0.15 g, 0.24 mmol) and 1-(O-tolyl)-piperazine (0.51 g, 2.90 mmol) at 25 °C. The reaction mixture was stirred for 24 h and diluted with EtOAc (100 mL) and H_2O (50 mL). The organic layer was separated, washed with H_2O (2 x 50 mL), dried over Na_2SO_4 , and concentrated. Column chromatography (silica, 5% MeOH/ CH_2Cl_2) afforded 1.08 g (88%) of the target compound, a white powder. TLC (silica, 10% MeOH/ CH_2Cl_2): $R_f = 0.38$. MS (electrospray): m/z 508.3 ($[\text{M}+\text{H}]^+$, $\text{C}_{28}\text{H}_{34}\text{ClN}_5\text{O}_2$ requires 507.2). ^1H NMR (CDCl_3 , 400 MHz, a mixture of two rotamers): 7.60 and 7.37 (AB pattern, $J_{ab} = 8.8$ Hz, 2H), 7.54 and 7.40 (AB pattern, $J_{ab} = 8.8$ Hz, 2H), 7.18-7.14 (m, 2H), 7.00-6.97 (m, 2H), 4.85 and 4.73 (AB pattern, $J_{ab} = 15.5$ Hz, 1H), 4.62 (s, 1H), 4.20-4.11 (m, 2H), 4.06-4.01 (m, 1H), 3.88-3.70 (m, 2H), 2.97-2.87 (m, 6H), 2.85-2.75 (m, 2H), 2.65-2.55 (m, 2H), 2.51-2.48 (m, 2H), 2.29 (s, 3H), 2.21 (s, 1.5H), 2.17 (s, 1.5H).

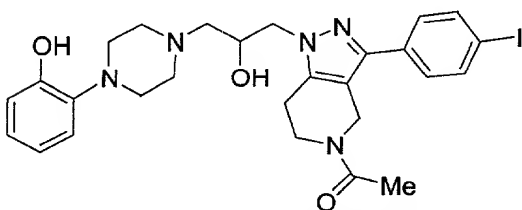
EXAMPLE 3



1-[3-(4-Chloro-phenyl)-1-[2-methoxy-3-(4-o-tolyl)-piperazin-1-yl]-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A stirred solution of 1-{3-(4-chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone (25 mg, 0.05 mmol) in THF (0.2 mL) was treated with NaH (1.42 mg, 0.06 mmol) at 25 °C. After 20 min, methyl iodide (3.7 µL, 0.06 mmol) was added and the reaction mixture was stirred for an additional 2 h. Preparative TLC (silica, 5% MeOH/CH₂Cl₂) afforded 14.6 mg (56%) of a colorless film. TLC (silica, 10% MeOH/CH₂Cl₂): R_f = 0.38. MS (electrospray): *m/z* 522.2 ([M+H]⁺, C₂₉H₃₆ClN₅O₂ requires 521.3). ¹H NMR (CDCl₃, 400 MHz, a mixture of two rotamers): 7.62 and 7.37 (AB pattern, *J*_{ab} = 8.8 Hz, 2H), 7.55 and 7.40 (AB pattern, *J*_{ab} = 8.8 Hz, 2H), 7.18-7.14 (m, 2H), 7.02-6.95 (m, 2H), 4.82 and 4.75 (AB pattern, *J*_{ab} = 15.5 Hz, 1H), 4.62 (s, 1H), 4.30-4.25 (m, 1H), 4.09-3.73 (m, 4H), 3.29 (s, 1.5H), 3.27 (s, 1.5H), 2.93-2.55 (m, 12H), 2.30 (s, 3H), 2.21 (s, 1.5H), 2.16 (s, 1.5H).

EXAMPLE 4



1-[1-{2-Hydroxy-3-[4-(2-hydroxy-phenyl)-piperazin-1-yl]-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A. 1-[3-(4-Iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A flask equipped with a Dean-Stark trap was charged with *N*-acetyl-4-piperidone (27.29g, 137 mmol), piperidine (16.5 mL, 129 mmol), *p*-toluenesulfonic acid (0.5 g) and benzene (150 mL). The mixture was heated to 125 °C. After 8 h the mixture was allowed to cool, and concentrated in vacuo to give the corresponding enamine (35 g). A solution of *p*-iodobenzoyl chloride (9.28 g, 34.8 mmol) in CH₂Cl₂ (40 mL) was added dropwise to a 0 °C solution of the enamine (11.0 g, ca. 41 mmol) in CH₂Cl₂ (80 mL) over 2 h. The mixture was then allowed to warm to room temperature and stirred for an additional 17 h. The solution was treated with 1 *N* HCl (200 mL) and stirred vigorously for 5

h. The layers were separated and the aqueous layer was extracted with CH_2Cl_2 (3 x 150 mL). The combined extracts were dried over Na_2SO_4 and concentrated. The residue was dissolved in EtOH (200 mL) and treated with NH_2NH_2 (16.0 mL, 51 mmol). The mixture was stirred for 17 h and H_2O (300 mL) was added. The precipitate formed was collected by filtration and air dried to give 8.82 g (59%) of 1-[3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone which was suitable for use without further purification. TLC (silica, 5% MeOH/ CH_2Cl_2): R_f = 0.3. MS (electrospray): m/z calculated for $\text{C}_{14}\text{H}_{15}\text{IN}_3\text{O}$ $[\text{M}+\text{H}]^+$ 368.03, found 368.0. ^1H NMR ($\text{CD}_3\text{OD}/\text{CDCl}_3$, 500 MHz, a mixture of amide rotamers): 7.72 (d, J = 8.2 Hz, 2H), 7.69 (d, J = 8.3 Hz, 2H), 7.24 (d, J = 8.2 Hz, 2H), 7.20 (d, J = 8.3 Hz, 2H), 4.69 (s, 2H), 4.56 (s, 2H), 3.83 (t, J = 6.0 Hz, 2H), 3.69 (t, J = 5.8 Hz, 2H), 2.79, (t, J = 5.7 Hz, 2H), 2.72, (t, J = 5.8 Hz, 2H), 2.13 (s, 3H), 2.08 (s, 3H).

B. 1-[3-(4-iodo-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

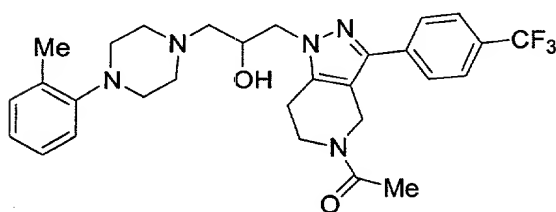
Cs_2CO_3 (1.30 g, 4.01 mmol) was added to a solution of 1-[3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (1.34 g, 3.65 mmol) and epichlorohydrin (2.85 mL, 36.4 mmol) in DMF (10.0 mL). The mixture was stirred for 17 h then partitioned between EtOAc (400 mL) and saturated NaHCO_3 (150 mL). The NaHCO_3 layer was extracted with EtOAc (2 x 150 mL). The combined extracts were washed with H_2O (2 x 150 mL), brine (150 mL), dried over Na_2SO_4 and concentrated. The residue was purified by column chromatography (silica, 10-25% acetone/ CH_2Cl_2) to give 890 mg (58%) of the title compound. HPLC, t_R = 5.53 min. (Reverse phase conditions: HP 1100 LCMS, Phenomenex luna 2.1 x 150 mm column, 60% MeOH/ H_2O (0.5% AcOH) to 90% MeOH/ H_2O (0.5% AcOH), held at initial conditions for 2 min then ramped to final conditions over 5 min.) MS (electrospray), m/z calculated for $\text{C}_{17}\text{H}_{18}\text{IN}_3\text{O}_2\text{Na}$ $[\text{M}+\text{Na}]^+$ 445.04, found 445.95. ^1H NMR (CDCl_3 , 500 MHz, a mixture of amide rotamers): 7.76 (d, J = 8.3 Hz, 2H), 7.75 (d, J = 8.3 Hz, 2H), 7.42 (d, J = 8.2 Hz, 2H), 7.35 (d, J = 8.2 Hz, 2H), 4.80 and 4.73 (A and B of AB quartet, J_{ab} = 15.6 Hz, 2H), 4.60 (s, 2H), 4.84 (dd, J = 15.1, 2.1 Hz, 1H), 4.42 (dd, J = 15.0, 2.1 Hz, 1H), 4.11 (t, J = 5.0 Hz, 1H), 4.08 (t, J = 5.0 Hz, 1H),

3.98-3.87 (m, 2H), 3.74 (m, 2H), 3.34 (m, 2H), 2.87-2.72 (m, 6H), 2.52 (dd, $J = 4.6, 2.6$ Hz, 1H), 2.48 (dd, $J = 4.5, 2.6$, 1H), 2.20 (s, 3H), 2.14 (s, 3H).

C. 1-[1-{2-Hydroxy-3-[4-(2-hydroxy-phenyl)-piperazin-1-yl]-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

1-[3-(4-Iodo-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (62 mg, 0.15 mmol) and 4-(2-hydroxyphenyl)-piperazine (34 mg, 0.19 mmol) were combined in CH_2Cl_2 (0.5 mL) and the solution treated with $\text{Yb}(\text{OTf})_3 \cdot \text{H}_2\text{O}$ (44 mg, 0.071 mmol). The mixture was shaken for 72 h then diluted with CH_2Cl_2 (1 mL). Purification by preparative TLC (silica, 10% MeOH/ CH_2Cl_2) gave 45 mg (51%) of an off-white powder. TLC (silica, 8% MeOH/ CH_2Cl_2): $R_f = 0.2$. MS (electrospray): m/z calculated for $\text{C}_{27}\text{H}_{33}\text{N}_5\text{O}_3$ $[\text{M}+\text{H}]^+$ 602.15, found 602.2. ^1H NMR (CDCl_3 , 500 MHz, a mixture of amide rotamers): 7.76 (d, $J = 8.6$ Hz, 1H), 7.72 (d, $J = 8.6$ Hz, 1H), 7.42 (d, $J = 8.5$ Hz, 1H), 7.34 (d, $J = 8.5$ Hz, 1H), 7.14 (m, 1H), 7.80 (t, $J = 7.7$ Hz, 1H), 6.94 (d, $J = 8.1$ Hz, 1H), 6.86 (t, $J = 7.7$ Hz, 1H), 4.83 and 4.72 (A and B of AB quartet, $J_{ab} = 15.6$ Hz, 1H), 4.61 (s, 1H), 4.22-4.15 (m, 2H), 4.02 (m, 2H), 3.88 (m, 1H), 3.76 (m, 3H), 3.00-2.49 (m, 11H), 2.20 (s, 1.5H), 2.15 (s, 1.5H).

EXAMPLE 5



1-[1-[2-Hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A. 1-[3-(4-Trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A solution of *N*-acetyl-4-piperidone (2.82 g, 20 mmol), morpholine (1.93 mL, 22 mmol) and *p*-toluenesulfonic acid (5 mg) in benzene (8.5 mL) was refluxed for

8 h in a Dean-Stark apparatus. The solvent was removed and the residue dissolved in CH_2Cl_2 (20 mL). Triethylamine (3.1 mL) was added and *p*-trifluoromethylbenzoyl chloride (3.27 mL, 22 mmol) in CH_2Cl_2 (4 mL) was added dropwise into the solution at 0 °C. The reaction mixture was stirred at 25 °C for 24 h and diluted with aqueous HCl (5%, 25 mL). After stirring for another 30 min, the organic layer was separated, washed with H_2O (20 mL), dried (Na_2SO_4), and concentrated. The residue was dissolved in EtOH (95%, 18 mL) and treated at 0 °C with hydrazine (2.9 mL, 60 mmol). The mixture was stirred at 25 °C for 3 h and H_2O (4 mL) was added. Most of the volatiles were removed and the residue extracted with CH_2Cl_2 (50 mL). The organic layer was separated, washed with H_2O (20 mL), dried over Na_2SO_4 , and concentrated. Column chromatography (silica, 5% MeOH/ CH_2Cl_2) provided 5.1 g (83%) of a white powder. TLC (silica, 10% MeOH/ CH_2Cl_2): R_f = 0.30. MS (electrospray): m/z 332.0 ($[\text{M}+\text{Na}]^+$, $\text{C}_{15}\text{H}_{14}\text{F}_3\text{N}_3\text{O}$ requires 309.1). ^1H NMR (CDCl_3 , 400 MHz, a mixture of two rotamers): 7.73-7.67 (m, 4H), 4.85 (s, 1.2H), 4.68 (s, 0.8H), 3.96 (t, J = 4.5 Hz, 0.8H), 3.78 (t, J = 4.5 Hz, 1.2H), 2.89 (t, J = 4.5 Hz, 1.2H), 2.83 (t, J = 4.5 Hz, 0.8H), 2.23 (s, 1.8H), 2.18 (s, 1.2H).

B. 1-[1-Oxiranylmethyl-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

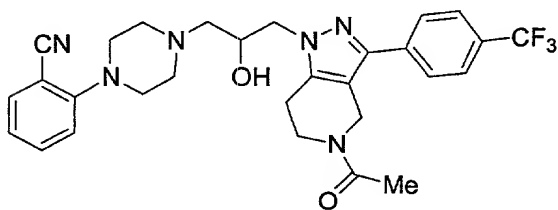
A solution of 1-[3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (2.4 g, 7.77 mmol) in DMF (15 mL) was treated with cesium carbonate (5.05 g, 15.5 mmol) and epichlorohydrin (6.1 mL, 77.7 mmol) at 25 °C and stirred for 24 h before it was diluted with EtOAc (100 mL) and H_2O (50 mL). The organic layer was separated, washed with H_2O (2 x 50 mL), brine (50 mL), dried over Na_2SO_4 , and concentrated. Column chromatography (silica, 10% acetone/ CH_2Cl_2) provided 2.30 g (81%) of a white powder. TLC (silica, 10% MeOH/ CH_2Cl_2): R_f = 0.35. MS (electrospray): m/z 388.0 ($[\text{M}+\text{Na}]^+$, $\text{C}_{18}\text{H}_{18}\text{F}_3\text{N}_3\text{O}_2$ requires 365.1). ^1H NMR (CDCl_3 , 400 MHz, a mixture of two rotamers): 7.77 and 7.63 (AB pattern, J_{ab} = 8.2 Hz, 2H), 7.71 and 7.67 (AB pattern, J_{ab} = 8.4 Hz, 2H), 4.82 and 4.76 (AB pattern, J_{ab} = 15.5 Hz, 1.2H), 4.58 (s, 0.8H), 4.45-4.35 (m, 1H), 4.08-4.02 (m, 1H), 3.92-3.80 (m, 1H), 3.70-3.63

(m, 1H), 3.30 (m, 1H), 2.80-2.67 (m, 3H), 2.48-2.42 (m, 1H), 2.13 (s, 1.3H), 2.08 (s, 1.7H).

C. 1-[1-[2-Hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A solution of 1-[1-oxiranylmethyl-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (1.16 g, 3.20 mmol) in CH₂Cl₂ (15 mL) was treated with ytterbium(III) triflate (0.40 g, 0.64 mmol) and 1-(O-tolyl)-piperazine (0.84 g, 4.77 mmol) at 25 °C and stirred for 48 h before it was diluted with CH₂Cl₂ (100 mL) and H₂O (50 mL). The organic layer was separated, washed with H₂O (2 x 50 mL), dried over Na₂SO₄, and concentrated. Column chromatography (silica, 5% MeOH/CH₂Cl₂) afforded 1.54 g (89%) of a white powder. TLC (silica, 10% MeOH/CH₂Cl₂): R_f = 0.35. MS (electrospray): *m/z* 542.3 ([M+H]⁺, C₂₉H₃₄F₃N₅O₂ requires 541.3). ¹H NMR (CDCl₃, 400 MHz, a mixture of two rotamers): 7.82 and 7.65 (AB pattern, *J*_{ab} = 8.2 Hz, 2H), 7.72 and 7.68 (AB pattern, *J*_{ab} = 8.4 Hz, 2H), 7.18-6.97 (m, 4H), 4.88 and 4.76 (AB pattern, *J*_{ab} = 16 Hz, 0.9H), 4.65 (s, 1.1H), 4.23-4.12 (m, 2H), 4.08-4.00 (m, 2H), 3.88-3.70 (m, 2H), 3.02-2.85 (m, 6H), 2.85-2.75 (m, 2H), 2.65-2.55 (m, 2H), 2.53-2.45 (m, 2H), 2.29 (s, 3H), 2.21 (s, 1.8H), 2.17 (s, 1.2H).

EXAMPLE 6



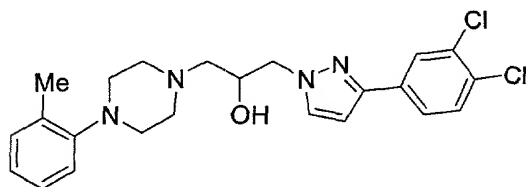
2-(4-{3-[5-Acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile.

A solution of 1-[1-oxiranylmethyl-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (0.84 g, 2.30 mmol) in CH₂Cl₂ (10 mL)

was treated with ytterbium(III) triflate (0.29 g, 0.46 mmol) and 1-(2-cyanophenyl)-piperazine (0.75 g, 3.5 mmol) at 25 °C and stirred for 48 h before it was diluted with CH₂Cl₂ (100 mL) and H₂O (50 mL). The organic layer was separated, washed with H₂O (2 x 50 mL), dried over Na₂SO₄, and

- 5 concentrated. Column chromatography (silica, 5% MeOH/CH₂Cl₂) afforded 1.15 g (90%) of light yellow crystals. TLC (silica, 10% MeOH/CH₂Cl₂): R_f = 0.30. MS (electrospray): *m/z* 553.3 ([M+H]⁺, C₂₉H₃₁F₃N₆O₂ requires 552.3). ¹H NMR (CDCl₃, 400 MHz, a mixture of two rotamers): 7.82 and 7.68 (AB pattern, *J*_{ab} = 8.2 Hz, 2H), 7.76 and 7.72 (AB pattern, *J*_{ab} = 8.4 Hz, 2H), 7.60-7.48 (m, 2H),
 10 7.05-7.00 (m, 2H), 4.90 and 4.78 (AB pattern, *J*_{ab} = 16 Hz, 1H), 4.69 (s, 1H), 4.30-3.71 (m, 6H), 3.25 (m, 4H), 3.02-2.75 (m, 4H), 2.70-2.65 (m, 2H), 2.60-2.53 (m, 2H), 2.23 (s, 1.5 H), 2.18 (s, 1.5H).

15 EXAMPLE 7



1-[3-(3,4-Dichloro-phenyl)-pyrazol-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol.

A. 3-(3,4-Dichloro-phenyl)-1-oxiranylmethyl-1H-pyrazole.

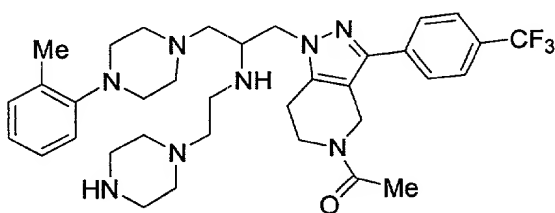
- 20 A stirred solution of 3-(3,4-dichlorophenyl)pyrazole (300 mg, 1.4 mmol) in DMF (5 mL) was treated with cesium carbonate (550 mg, 1.69 mmol) and epichlorohydrin (1.1 mL, 14.0 mmol) at room temperature for 18 h. The crude reaction mixture was then partitioned between EtOAc (50 mL) and water (35 mL). The aqueous phase was further extracted (2 x 50 mL) and the combined
 25 organic layers were washed with brine, dried (Na₂SO₄), filtered and concentrated under reduced pressure to yield crude product. Purification by column chromatography (silica, 25% EtOAc/hexanes) afforded 308 mg (82%) of the title compound. ¹NMR (400 MHz, CDCl₃): 7.83 (d, *J* = 2 Hz, 1H), 7.54 (dd, *J* = 2, 8 Hz, 1H), 7.44 (d, *J* = 2 Hz, 1H), 7.38 (d, *J* = 8 Hz, 1H), 6.48 (d, *J* = 2

Hz, 1H), 4.45 (dd, $J = 3, 9.7$ Hz, 1H), 4.12 (dd, $J = 6, 15$ Hz, 1H), 3.31 (m, 1H), 2.81 (dd, $J = 4.0, 4.6$ Hz, 1H), 2.47 (dd, $J = 2.6, 4.7$ Hz, 1H).

B. 1-[3-(3,4-Dichloro-phenyl)-pyrazol-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol.

A solution of 3-(3,4-dichloro-phenyl)-1-oxiranylmethyl-1H-pyrazole (30 mg, 0.11 mmol) and 1-(2-methylphenyl)-piperazine (22 mg, 0.12 mmol) in EtOH (1 mL) was heated to 80 °C overnight. Removal of solvent and purification by column chromatography (silica, 0-5% acetone/ CH_2Cl_2) afforded 35 mg (70%) of the title compound. ^1H NMR (400 MHz, CDCl_3): 7.89 (d, $J = 2$ Hz, 1H), 7.61 (dd, $J = 2, 8.7$ Hz, 1H), 7.57 (d, $J = 2$ Hz, 1H), 7.45 (d, $J = 8.4$ Hz, 1H), 7.16 (m, 2H), 6.99 (m, 2H), 6.54 (d, $J = 2.3$ Hz, 1H), 4.31 (m, 1H), 4.18 (m, 2H), 2.93 (m, 4H), 2.60 (m, 2H), 2.47 (m, 3H), 2.88 (s, 3H).

EXAMPLE 8



1-[1-[2-(2-Piperazin-1-yl-ethylamino)-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A. 1-[5-Acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-one.

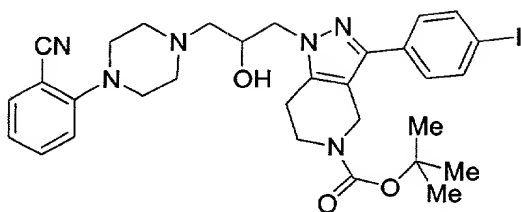
A solution of DMSO (3.55 mL, 50 mmol) in CH_2Cl_2 (7 mL) was treated with oxalyl chloride (2.90 mL, 33 mmol) at -78 °C and stirred for 30 min. A solution of 1-[1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (1.8 g, 3.3 mmol) in CH_2Cl_2 (7 mL) was then slowly added and the reaction mixture was stirred for an additional 30 min before it was quenched with addition of triethylamine (18.4 mL, 132 mmol). The reaction mixture was slowly warmed to

25 °C and diluted with EtOAc (50 mL) and sat. NaHCO₃ (30 mL). The organic layer was separated, washed with H₂O (2 x 50 mL), dried over Na₂SO₄, and concentrated. Column chromatography (silica, 2-5% MeOH/CH₂Cl₂) afforded 1.50 g (83%) of a light yellow powder. TLC (silica, 10% MeOH/CH₂Cl₂): R_f = 0.35. MS (electrospray): *m/z* 540.3 ([M+H]⁺, C₂₉H₃₂F₃N₅O₂ requires 539.3). ¹H NMR (CDCl₃, 400 MHz, a mixture of two rotamers): 7.78 and 7.62 (AB pattern, J_{ab} = 8.2 Hz, 2H), 7.70 and 7.64 (AB pattern, J_{ab} = 8.4 Hz, 2H), 7.18-6.95 (m, 4H), 5.10 (s, 1H), 5.07 (s, 1H), 4.84 (s, 1H), 4.68 (s, 1H), 3.96 (t, J = 4.4 Hz, 1H), 3.78 (t, J = 4.4 Hz, 1H), 3.47 (s, 4H), 3.34 (s, 2H), 2.74-2.65 (m, 6H), 2.29 (s, 3H), 2.20 (s, 1.5H), 2.17 (s, 1.5H).

B. 1-[1-[2-(2-Piperazin-1-yl-ethylamino)-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A solution of 1-[5-acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-one (54 mg, 0.1 mmol) in 1,2-dichloroethane (0.5 mL) was treated with 1-(2-aminoethyl)piperazine (26 µL, 0.2 mmol) and glacial acetic acid (34 µL, 0.6 mmol) at 25 °C and stirred for 30 min. Sodium triacetoxyborohydride (63.6 mg, 0.3 mmol) was added and the reaction mixture was stirred for an additional 4 h before it was quenched with CH₂Cl₂ (5 mL) and sat. NaHCO₃ (5 mL). The organic layer was separated, washed with H₂O (2 x 5 mL), dried over Na₂SO₄, and concentrated. Preparative TLC (silica, 10% MeOH/CH₂Cl₂) afforded 22 mg (35%) of a light yellow film. TLC (10% MeOH/CH₂Cl₂): R_f = 0.2. MS (electrospray): *m/z* 653.3 ([M+H]⁺, C₃₅H₄₇F₃N₈O requires 652.4). ¹H NMR (CDCl₃, 400 MHz, a mixture of two rotamers): 7.78-7.60 (m, 4H), 7.18-6.82 (m, 4H), 4.88-4.30 (m, 2H), 4.23-3.90 (m, 2H), 3.85-3.70 (m, 2H), 3.22-2.85 (m, 10H), 2.85-2.30 (m, 15H), 2.30 (s, 3H), 2.21 (s, 1.5H), 2.17 (s, 1.5H).

EXAMPLE 9



- 5 1-{3-[4-(2-Cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

A. 3-(4-Iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

- 10 *p*-Toluenesulfonic acid (0.055 g, 0.29 mmol) and morpholine (4.76 mL, 54 mmol) were added to a solution of *tert*-butyl 4-oxo-1-piperidinecarboxylate (10.3 g, 52 mmol) in benzene (22 mL). The reaction mixture was heated in a flask equipped with a condenser and a Dean-Stark trap at reflux for 20 h. The reaction mixture was cooled and concentrated in vacuo to give the enamine
- 15 which was used without further purification. The enamine was dissolved in CH₂Cl₂ (60 mL) and cooled to 0 °C. Triethylamine (8.67 mL, 62 mmol) was added, followed by dropwise addition of 4-iodobenzoyl chloride (13.8 g, 52 mmol) dissolved in CH₂Cl₂ (10 mL). The reaction mixture was allowed to warm to room temperature and stirred for 72 h. The reaction mixture was poured
- 20 over water (200 mL) and the CH₂Cl₂ layer was separated, dried (Na₂SO₄), and concentrated. The resulting oil was taken up in EtOH (200 mL) and treated with hydrazine (4.88 mL, 155 mmol) at 0 °C. The reaction mixture was allowed to warm to room temperature and stirred for 17 h. The mixture was concentrated and the resulting material was triturated with EtOAc to afford 9.52
- 25 g (43%) of a white solid. TLC (silica, 10% acetone/CH₂Cl₂): *R*_f = 0.18. MS (electrospray): *m/z* 426.0 (426.1 calculated for C₁₇H₂₀IN₃O₂, [M+H]⁺). ¹H NMR (400 MHz, CDCl₃): 7.74 (br s, 2H), 7.31 (br d, *J* = 8.0 Hz, 2H), 4.63 (br s, 2H), 3.73 (br s, 2H), 2.77 (br s, 2H), 1.49 (s, 9H).

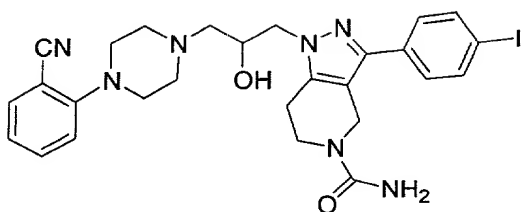
- 30 B. 3-(4-Iodo-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

Cesium carbonate (1.84 g, 5.65 mmol) was added to a solution of epichlorohydrin (3.68 mL, 47.05 mmol) and 3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (2.0 g, 4.71 mmol) in DMF (10 mL). The reaction mixture was allowed to stir for 24 h, then partitioned between aqueous NaHCO₃ and EtOAc. The aqueous layer was extracted with EtOAc and the combined organic layers were washed with water and brine, dried (Na₂SO₄), and concentrated. Purification by column chromatography (silica, 0–10% acetone/CH₂Cl₂) afforded 2.26 g (69%) of a white foam. TLC (silica, 10% acetone/CH₂Cl₂): *R*_f = 0.44. MS (electrospray): *m/z* 482.0 (482.1 calculated for C₂₀H₂₄IN₃O₃, [M+H]⁺). ¹H NMR (400 MHz, CDCl₃): 7.60 (br s, 2H), 7.28 (br d, *J* = 8.2 Hz, 2H), 4.48 (br s, 2H), 4.32 (br d, *J* = 14.8 Hz, 1H), 3.99 (dd, *J* = 15.0, 5.4 Hz, 1H), 3.61 (br s, 1H), 3.26–3.20 (m, 1H), 2.72 (t, *J* = 4.4 Hz, 1H), 2.65–2.58 (m, 2H), 2.40 (br s, 1H), 1.36 (s, 9H).

C. 1-{3-[4-(2-Cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

Ytterbium (III) trifluoromethanesulfonate hydrate (0.193 g, 0.311 mmol) and 1-(2-cyanophenyl)-piperazine (0.292 g, 1.56 mmol) were dissolved in CH₂Cl₂ (2 mL) and added to a solution of 3-(4-iodo-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester in CH₂Cl₂ (5 mL). The reaction mixture was allowed to stir for 48 h at 25 °C. Purification by flash chromatography (silica, 0–15% acetone/CH₂Cl₂) afforded 392 mg (56%) of a white foam. TLC (silica, 10% acetone/CH₂Cl₂): *R*_f = 0.41. MS (electrospray): *m/z* 669.2 (669.2 calculated for C₃₁H₃₇IN₆O₃, [M+H]⁺). ¹H NMR (400 MHz, CDCl₃): 7.73 (br s, 2H), 7.58–7.56 (m, 1H), 7.52–7.48 (m, 1H), 7.39 (br d, *J* = 7.1 Hz, 2H), 7.04–7.00 (m, 2H), 4.60 (br s, 2H), 4.06–4.04 (m, 2H), 4.06–4.04 (m, 1H), 3.76–3.70 (m, 2H), 3.26 (br s, 4H), 2.84–2.38 (m, 7H), 1.56–1.53 (m, 2H), 1.48 (s, 9H).

EXAMPLE 10



1-{3-[4-(2-Cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-iodo-phenyl)-
 1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide.

A. 2-(4-{2-Hydroxy-3-[3-(4-iodo-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzonitrile.

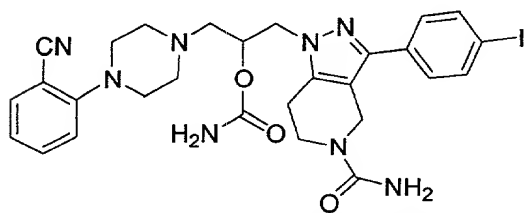
Trifluoroacetic acid (3 mL) was added to a solution of 1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (0.402 g, 0.601 mmol) in CH_2Cl_2 (3 mL) and the reaction mixture was stirred for 2 h. The mixture was concentrated, then diluted with EtOAc. The organic layer was washed with aqueous NaHCO_3 and brine, dried (Na_2SO_4), and concentrated to afford the amine (0.342 g, 100%) as a yellowish foam. TLC (silica, 10% acetone/ CH_2Cl_2): R_f = 0.14. MS (electrospray): m/z 569.2 (569.1, calculated for $\text{C}_{26}\text{H}_{29}\text{IN}_6\text{O}$, $[\text{M}+\text{H}]^+$). ^1H NMR (400 MHz, CDCl_3 : CD_3OD (6:1)): 7.73 (d, J = 8.6 Hz, 2H), 7.56 (dd, J = 7.6, 1.8 Hz, 1H), 7.52 (t, J = 8.0 Hz, 1H), 7.25 (d, J = 8.6 Hz, 2H), 7.09 (t, J = 7.6 Hz, 1H), 7.02 (dd, J = 8.4 Hz, 1H), 4.43-4.36 (m, 1H), 4.31 (s, 2H), 4.21 (dd, J = 14.1, 4.5 Hz, 1H), 4.11 (dd, J = 14.5, 6.3 Hz, 1H), 3.54-3.49 (m, 2H), 3.40-3.24 (m, 8H), 3.18-3.06 (m, 3H), 3.03-2.95 (m, 3H).

B. 1-{3-[4-(2-Cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide.

Diisopropylethylamine (0.531 mL, 3.05 mmol), DMAP (5 mg), and trimethylsilyl isocyanate (0.413 mL, 3.05 mmol) were added to a solution of 2-(4-{2-hydroxy-3-[3-(4-iodo-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzonitrile in pyridine (3 mL) and CH_2Cl_2 (6 mL). The reaction mixture was stirred for 20 h, then partitioned between aqueous NaHCO_3 and CH_2Cl_2 . The CH_2Cl_2 layer was washed with brine, dried (Na_2SO_4), and

concentrated. The resulting product was dissolved in CH_2Cl_2 (5 mL) and treated with 21 wt% sodium ethoxide in EtOH (0.5 mL) for 3 h. The reaction mixture was washed with brine, dried (Na_2SO_4), and concentrated. Purification by column chromatography (silica, 0-10% MeOH/ CH_2Cl_2) afforded 290 mg (78%) of the title compound. HPLC (reverse phase conditions), $t_R = 4.21$ min. MS (electrospray): m/z 612.2 (612.5, calculated for $\text{C}_{27}\text{H}_{30}\text{IN}_7\text{O}_2$, $\text{M}^+ + \text{H}$). ^1H NMR (400 MHz, CDCl_3): 7.73 (d, $J = 8.6$ Hz, 2H), 7.57 (dd, $J = 7.6$, 1.6 Hz, 1H), 7.50 (t, $J = 7.8$ Hz, 1H), 7.36 (d, $J = 8.6$ Hz, 2H), 7.05 (t, $J = 7.6$ Hz, 1H), 7.00 (d, $J = 8.4$ Hz, 1H), 4.64 (br s, 2H), 4.57 (br s, 2H), 4.30 (br s, 1H), 4.20 (dd, $J = 14.1$, 3.3 Hz, 1H), 4.06 (dd, $J = 14.1$, 6.3 Hz, 1H), 3.82-3.65 (m, 2H), 3.29-3.20 (m, 4H), 3.04-2.80 (m, 6H), 2.68 (br s, 2H).

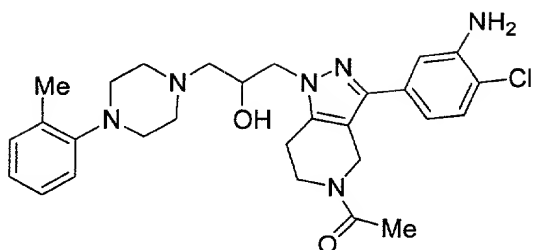
EXAMPLE 11



Carbamic acid 1-[5-carbamoyl-3-(4-iodo-phenyl)-4,5,6,7-tetrahydropyrazolo[4,3-c]pyridin-1-ylmethyl]-2-[4-(2-cyano-phenyl)-piperazin-1-yl]-ethyl ester.

The title compound (13 mg, 3%) was obtained along with 1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydropyrazolo[4,3-c]pyridine-5-carboxylic acid as described in example 10. MS (electrospray): m/z 655.2 (655.2, calculated for $\text{C}_{28}\text{H}_{31}\text{IN}_8\text{O}_3$, $[\text{M} + \text{H}]^+$). HPLC (reverse phase conditions): $t_R = 6.29$ min. ^1H NMR (400 MHz, CDCl_3): 7.69 (d, $J = 8.1$ Hz, 2H), 7.50 (d, $J = 7.52$, 1H), 7.43 (t, $J = 8.1$ Hz, 1H), 7.31 (d, $J = 8.1$ Hz, 2H), 6.96 (t, $J = 9.0$ Hz, 2H), 4.64 (br s, 2H), 4.08 (d, $J = 16.8$ Hz, 2H), 3.96 (dd, $J = 14.0$, 6.6 Hz, 1H), 3.80-3.69 (m, 2H), 3.10-2.80 (m, 4H), 2.66 (br s, 2H), 2.50 (br s, 2H).

EXAMPLE 12



1-{3-(3-Amino-4-chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-
5 1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone.

A. 1-[3-(4-Chloro-3-nitro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A flask equipped with a Dean-Stark trap was charged with *N*-acetyl-4-
10 piperidone (27.29g, 137 mmol), piperidine (16.5 mL, 129 mmol), *p*-toluene-sulfonic acid (0.5 g) and benzene (150 mL). The mixture was heated to 125 °C. After 8 h the mixture was allowed to cool, and concentrated in vacuo to give the corresponding enamine (35 g). A solution of the enamine (3.87 g, 20.0 mmol) in dichloromethane (24 mL) was treated with triethylamine (3.07
15 mL, 22.0 mmol) and 4-chloro-3-nitrobenzoyl chloride (4.84 g, 22.0 mmol). The reaction mixture was stirred at 0 °C for 1 h and then at room temperature for 16 h. Hydrazine (1.88 mL, 60 mmol) was added to the reaction mixture. This solution was stirred at room temperature for an additional 16 h. The solvents were removed under reduced pressure. Ethyl acetate (100 mL) was added to
20 the residue to form a suspension. This suspension was filtered and dried to afford 6.4 g (100%) of a yellow solid. MS (electrospray): *m/z* 321.0 (321.0, calculated for C₁₄H₁₃ClN₄O₃, [M+H]⁺). ¹H NMR (CDCl₃, 400 MHz, a mixture of two rotamers): 8.10-8.00 (m, 3H), 4.90 (s, 0.8H), 4.85 (s, 1.2H), 3.96 (m, 2H), 2.95 (m, 2H), 2.20 (s, 3H).

B. 1-[3-(4-Chloro-3-nitro-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A solution of 1-[3-(4-chloro-3-nitro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (6.4 g, 20.0 mmol) in DMF (60 mL) was treated with
30 cesium carbonate (13.0 g, 40 mmol) and epichlorohydrin (15.6 mL, 200.0

mmol) at room temperature. The reaction mixture was stirred at room temperature for an additional 24 h before it was diluted with ethyl acetate (350 mL) and water (50 mL). The organic layer was separated, washed with water (2 x 50 mL), brine (50 mL), dried over sodium sulfate, and concentrated under reduced pressure. The residue was purified by column chromatography (silica, 10% acetone/CH₂Cl₂) to provide 7.5 g (83%) of a light yellow powder. MS (electrospray): *m/z* 377.0 (377.0, calculated for C₁₇H₁₇ClN₄O₄, [M+H]⁺). ¹H NMR (CDCl₃, 400 MHz, a mixture of two rotamers): 8.15-8.05 (m, 1H), 7.75-7.65 (m, 1H), 7.55-7.45 (m, 1H), 4.80-4.65 (m, 1.2H), 4.60 (s, 0.8H), 4.45-4.35 (m, 1H), 4.08-4.02 (m, 1H), 3.92-3.80 (m, 1H), 3.70-3.63 (m, 1H), 3.30-3.20 (m, 1H), 2.90-2.67 (m, 3H), 2.55-2.48 (m, 1H), 2.15 (s, 1.7H), 2.10 (s, 1.3H).

C. 1-[3-(4-Chloro-3-nitro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A solution of 1-[3-(4-chloro-3-nitro-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (0.754 g, 2.0 mmol) in dichloromethane (10 mL) was treated with ytterbium(III) triflate (0.25 g, 0.40 mmol) and 1-(2-methylphenyl)-piperazine (0.705 g, 4.0 mmol) at room temperature. The reaction mixture was stirred at room temperature for 16 h and diluted with dichloromethane (100 mL) and water (50 mL). The organic layer was separated, washed with water (2 x 50 mL), dried over sodium sulfate and concentrated under reduced pressure. The residue was purified by column chromatography (silica, 5% MeOH/CH₂Cl₂) to afford 0.98 g (90%) of the desired product as a light yellow solid. MS (electrospray): *m/z* 553.2 (553.2, calculated for C₂₈H₃₃ClN₆O₄, [M+H]⁺). ¹H NMR (CDCl₃, 400 MHz, a mixture of two rotamers): 8.25-8.15 (m, 1H), 7.75-7.70 (m, 1H), 7.63-7.55 (m, 1H) 7.20-7.10(m, 2H), 7.05-6.95 (m, 2H), 4.90-4.70 (m, 1H), 4.65 (s, 1H), 4.30-4.15 (m, 2H), 4.10-3.70 (m, 4H), 3.00-2.40 (m, 12H), 2.20(s, 3H), 2.15 (s, 1.5H), 2.10 (s, 1.5H).

D. 1-[3-(3-amino-4-chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

To a solution of sodium hydrosulfite (1.28 g, 7.3 mmol) in 30 mL water was added 1-{3-(4-chloro-3-nitro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone (810 mg, 1.5 mmol) in 15 mL THF. The reaction mixture was stirred at room temperature for

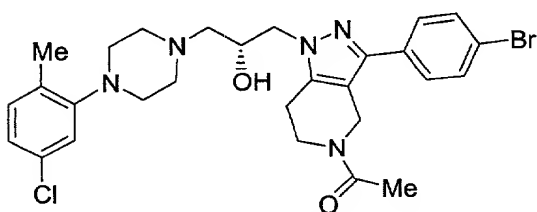
5 min. The color of the solution changed from light yellow to colorless.

Hydrochloride solution (1 N, 10 mL) was added to the reaction mixture. This solution was stirred at room temperature for 30 min, and treated with saturated sodium bicarbonate until the pH of the solution between 7 to 8. The product was extracted with dichloromethane (3 x 80 mL). The organic phases were

combined, dried over sodium sulfate, and concentrated under reduced pressure to a residue. This residue was purified by column chromatography (silica, 5-20% MeOH/CH₂Cl₂) to afford 644 mg (84.1%) of the title compound.

MS (electrospray): *m/z* 523.3 (523.3, calculated for C₂₈H₃₅ClN₆O₂, [M+H]⁺). ¹H NMR (CDCl₃, 400 MHz, a mixture of two rotamers): 7.30-6.70 (m, 7H), 4.80-4.60 (m, 1H), 4.55 (s, 1H), 4.20-4.05 (m, 4H), 3.95-3.90 (m, 2H), 3.80-3.60 (m, 2H), 2.90-2.30 (m, 9H), 2.20 (s, 3H), 2.15 (s, 1.5H), 2.10 (s, 1.5H).

EXAMPLE 13



(*R*)-1-(3-(4-Bromo-phenyl)-1-{3-[4-(5-chloro-2-methyl-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone.

A. (2*S*)-1-*tert*-Butyldimethylsilylglycidol.

tert-Butylchlorodimethylsilane (9.41 g, 62.4 mmol) followed by Et₃N (13.5 mL, 96.8 mmol) was added to a 0 °C solution of *R*-(+)-glycidol (3.88 g, 52.4 mmol) in CH₂Cl₂ (100 mL). The solution was allowed to warm to 23 °C with stirring over 17h. The resulting pink solution was diluted with Et₂O (250 mL) and stirred an additional 30 min. The solution was partitioned between Et₂O (800

mL) and sat. aqueous NaHCO₃ (200 mL). The Et₂O layer was washed with sat. aqueous NaHCO₃ (250 mL), H₂O (3 x 200 mL), brine (100 mL), dried over Na₂SO₄ and concentrated. Purification of the residue by column chromatography (silica, 5-10% Et₂O/hexanes) provided 8.21 g (84%) of the title compound. TLC (silica, 10% Et₂O/hexanes): R_f = 0.5. ¹H NMR (CDCl₃, 400 MHz): 3.85 (dd, *J* = 11.9, 3.2 Hz, 1H), 3.66 (dd, *J* = 11.9, 4.8 Hz, 1H), 3.09 (m, 1H), 2.77 (dd, *J* = 5.0, 4.2 Hz, 1H), 2.64 (dd, *J* = 5.2, 2.7 Hz, 1H), 0.90 (s, 9H), 0.08 (s, 3H), 0.07 (s, 3H).

10 B. 1-[3-(4-Bromo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A flask equipped with a Dean-Stark trap was charged with *N*-acetyl-4-piperidone (100.1 g, 709 mmol), piperidine (68 mL, 779 mmol), *p*TsOH (3.7 g) and benzene (500 mL). The mixture was heated to 125 °C. After 17 h the mixture was allowed to cool and divided into two portions. A solution of *p*-bromobenzoyl chloride (70.0 g, 319 mmol) in CH₂Cl₂ (400 mL) was added dropwise to a 0 °C solution of the enamine (ca. 355 mmol) in CH₂Cl₂ (320 mL) over 15 h. The mixture was then allowed to warm to 23 °C and stirred for an additional 5 h. The solution was treated with 1 *N* HCl (500 mL) and stirred vigorously for 1.5 h. The layers were separated and the aqueous layer was extracted with CH₂Cl₂ (2 x 300 mL). The combined extracts were washed with sat. aqueous NaHCO₃ (300 mL), H₂O (300 mL), brine (300 mL), dried over Na₂SO₄ and concentrated. The residue was dissolved in MeOH (300 mL) and treated with NH₂NH₂ (50.0 mL, 1.59 mol). The mixture was stirred for 17 h before the precipitate formed was collected by filtration and air dried to give 52 g (50%) of the title compound which was suitable for use without further purification. TLC (silica, 5% MeOH/CH₂Cl₂): R_f = 0.3. MS (electrospray): *m/z* calculated for C₁₄H₁₅⁷⁹BrN₃O [M+H]⁺, 320.04, found 320. ¹H NMR (CD₃OD/CDCl₃, 400 MHz, a mixture of amide rotamers): 7.53 and 7.35 (A and B of AA'BB', *J* = 8.5 Hz, 2H), 7.51 and 7.39 (A and B of AA'BB', *J* = 8.6 Hz, 2H), 4.72 (s, 2H), 4.58 (s, 2H), 3.85 (t, *J* = 5.9 Hz, 2H), 3.71 (t, *J* = 5.8 Hz, 2H), 2.81, (t, *J* = 5.8 Hz, 2H), 2.74, (t, *J* = 5.8 Hz, 2H), 2.16 (s, 3H), 2.11 (s, 3H).

C. (S)-1-[3-(4-Bromo-phenyl)-1-oxiranylmethyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

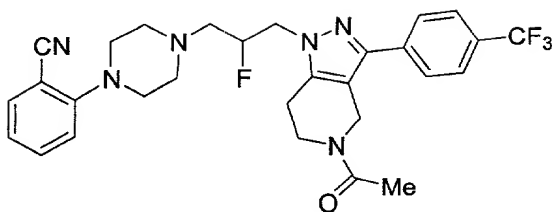
A solution of KHMDS in toluene (0.5 M, 3.7 mL, 1.85 mmol) was added to a 0 °C solution of 1-[3-(4-bromo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (492 mg, 1.54 mmol) in DMF (4.0 mL). The mixture was stirred for 1 h before (2S)-1-*tert*-butyldimethylsilylglycidol (870 mg, 4.62 mmol) was added neat via syringe. The mixture was stirred an additional 48 h and partitioned between EtOAc (300 mL) and sat. aqueous NaHCO₃ (100 mL). The aqueous layer was extracted with EtOAc (2 x 100 mL). The combined extracts were washed with H₂O (2 x 100 mL), brine (100 mL), dried over Na₂SO₄ and concentrated. The residue was dissolved in MeOH (50 mL) and treated with CSA (97 mg). The mixture was stirred for 17 h and concentrated to dryness. The residue was suspended in MeC(OMe)₃ (50 mL) and stirred for an additional 17 h. The mixture was diluted with EtOAc (400 mL) and washed with saturated aqueous NaHCO₃ (50 mL), H₂O (2 x 50 mL), brine (50 mL), dried over Na₂SO₄ and concentrated. The crude orthoester was dissolved in CH₂Cl₂ (5 mL), cooled to 0 °C, and treated with AcBr (0.18 mL, 2.4 mmol). The mixture was allowed to warm with stirring over 4 h before being worked up as described above. The crude acetyl-bromide obtained was dissolved in MeOH (50 mL), treated with K₂CO₃ (207 mg, 1.50 mmol) and stirred for 4 h. The reaction mixture was diluted with EtOAc (400 mL) and washed with saturated aqueous NH₄Cl (100 mL). The EtOAc layer was washed with H₂O (2 x 100 mL), brine (100 mL), dried over Na₂SO₄ and concentrated. The crude product was purified by column chromatography (silica, 10-40% acetone/CH₂Cl₂) to afford 158 mg (27%) of the title compound. Chiral HPLC (Daicel OD, 0.5% Et₂NH/MeOH) analysis indicated >95% optical purity. HPLC (reverse phase conditions): *t*_R = 4.90 min. MS (electrospray): *m/z* calculated for C₁₇H₁₉⁷⁹BrN₃O₂ [M⁺+H], 376.07, found 376.0. ¹H NMR (CDCl₃, 400 MHz, a mixture of amide rotamers): 7.47 (d with fine splittings (partially obscured), *J* = 8.5, Hz, 2H), 7.44 (m, 4H), 7.38 (d with fine splittings, *J* = 8.5, Hz, 2H), 4.71 and 4.64 (A and B of AB quartet, *J*_{ab} = 15.7 Hz, 2H), 4.51 (s, 2H), 4.39 (dd, *J* = 15.1, 2.5 Hz, 1H), 4.34 (dd, *J* = 15.0, 2.9 Hz, 1H), 4.02 (dd, *J* = 5.2, 3.9 Hz, 1H), 3.98 (dd, *J* = 5.3, 3.7 Hz, 1H), 3.83 (m, 2H), 3.64 (m, 2H), 3.25 (br m, 2H), 2.80-2.60 (m, 6H),

2.46 (dd, $J = 4.6, 2.6$ Hz, 1H), 2.38 (dd, $J = 4.6, 2.6$ Hz, 1H), 2.10 (s, 3H), 2.06 (s, 3H).

D. (R)-1-(3-(4-Bromo-phenyl)-1-{3-[4-(5-chloro-2-methyl-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone.

(S)-1-[3-(4-Bromo-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone (37 mg, 0.98 mmol) and 4-(2-methyl-5-chlorophenyl)piperazine (36 mg, 0.17 mmol) were combined in EtOH (0.4 mL) and heated to 70 °C. After 18 h the mixture was allowed to cool, diluted with CH₂Cl₂ and purified by preparative TLC (silica, 8% MeOH/CH₂Cl₂) to give 35 mg (61%) the title compound. HPLC (reverse phase conditions): $t_R = 4.41$ min. MS (electrospray): m/z calculated for C₂₈H₃₄³⁵Cl⁷⁹BrN₅O₂ [M⁺+H], 586.16, found 586.2. ¹H NMR (CDCl₃, 400 MHz, a mixture of amide rotamers): 7.56 (d (partially obscured), $J = 8.5$, Hz, 2H), 7.53 (s, 4H), 7.48 (d, $J = 8.5$ Hz, 2H), 7.08 (br d, $J = 8.5$ Hz, 1H), 6.95 (m, 2H), 4.85 and 4.73 (A and B of AB quartet, $J_{ab} = 15.6$ Hz, 1H), 4.62 (s, 1H), 4.20 (m, 2H), 4.04 (m, 2H), 3.90-3.71 (m, 2H), 2.92-2.53 (m, 11H), 2.21 (s, 1.5H), 2.16 (s, 1.5H).

EXAMPLE 14

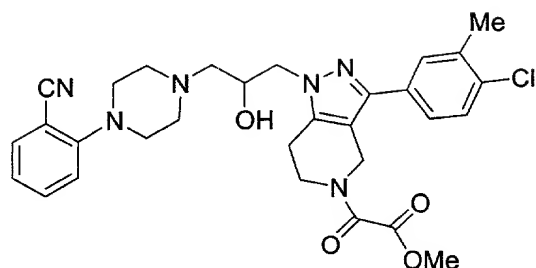


- 5 2-(4-{3-[5-Acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-fluoro-propyl}-piperazin-1-yl)-benzonitrile.

A solution of 2-(4-{3-[5-acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile (150
 10 mg, 0.27 mmol) in CH_2Cl_2 (1 mL) was treated with DAST (Et_2NSF_3 , 7 μL , 0.60 mmol) at -78°C . The reaction mixture was slowly warmed to 25°C for 1 h and then to 60°C for an additional 2 h. Preparative TLC (silica, 5% $\text{MeOH}/\text{CH}_2\text{Cl}_2$) provided 75 mg (50%) of the title compound as a light yellow powder. TLC (5% $\text{MeOH}/\text{CH}_2\text{Cl}_2$): $R_f = 0.28$. MS (electrospray): m/z 555.2 ($[\text{M}+\text{H}]^+$, $\text{C}_{29}\text{H}_{30}\text{F}_4\text{N}_6\text{O}$ requires 554.2). ^1H NMR (CDCl_3 , 400 MHz, a mixture of two rotamers): 7.71 and 7.59 (AB pattern, $J_{ab} = 8.2$ Hz, 2H), 7.66 and 7.62 (AB pattern, $J_{ab} = 8.4$ Hz, 2H), 7.50-7.38 (m, 2H), 6.96-6.92 (m, 2H), 5.01 (dp, $J = 49.0$, 3.0 Hz, 1H), 4.77 and 4.73 (AB pattern, $J_{ab} = 15.7$ Hz, 1.1H), 4.59 (s, 0.9H), 4.41-4.18 (m, 2H), 3.95-3.80 (m, 1H), 3.69 (dd, $J = 5.5$, 5.5 Hz, 1H), 3.18 (m, 4H), 2.83-2.65
 15 (m, 8H), 2.14 (s, 1.6H), 2.10 (s, 1.4H).

20

EXAMPLE 15



(3-(4-Chloro-3-methyl-phenyl)-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-oxo-acetic acid methyl ester.

A. 4-Chloro-3-methyl-benzoyl chloride.

To a suspension of 52.55 g (0.31 mol) of 4-chloro-3-methyl-benzoic acid in CH_2Cl_2 (1.2 L) with DMF (1 mL) at 0 °C under N_2 with an outlet sparging through 2.5 *N* sodium hydroxide was added 29.56 mL (0.339 mol) of oxalyl chloride. The mixture was allowed to warm to room temperature over a 3 h period. The reaction mixture was concentrated and taken forward crude.

B. 3-(4-Chloro-3-methyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

To a stirred solution of 55.8 g (0.28 mol) of 4-oxo-piperidine-1-carboxylic acid tert-butyl ester and 25.7 g (0.29 mol) of morpholine in benzene (125 mL) was added a catalytic amount (~0.25 g) of *p*-toluenesulfonic acid. The mixture was heated to reflux for 10 h under a Dean-Stark trap. The solvent was removed under reduced pressure to give a brown oil. The crude product was diluted with CH_2Cl_2 (400 mL), and 46.83 mL (0.34 mol) of Et_3N was added. The mixture was cooled to 0 °C, and a solution of 4-chloro-3-methyl-benzoyl chloride (0.35 mol) in CH_2Cl_2 (200 mL) was added slowly by dropping funnel over 2 h. The reaction mixture was poured over water (400 mL) and the CH_2Cl_2 layer was separated, dried (Na_2SO_4), and concentrated. The resulting oil was taken up in EtOH (400 mL) and treated with 35 mL of hydrazine at 0 °C. The reaction mixture was allowed to warm to room temperature and stirred for 17 h, during which time a white precipitate formed. The volume of the reaction mixture was reduced to ~150 mL, and Et_2O (750 mL) was added. The

suspension was stirred vigorously for 2 h and was filtered then washed with Et₂O (2 x 200 mL) and dried under vacuum to afford 50.74 g (52% over 3 steps) of 3-(4-chloro-3-methyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester as a pale orange solid. MS

- 5 (electrospray): exact mass calculated for C₁₈H₂₂ClN₃O₂, 347.1; *m/z* found, 348.1 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃): 7.26-7.43 (m, 4H), 4.65 (br s, 2H), 3.73 (br s, 2H), 2.77 (br s, 2H), 2.34 (s, 3H), 1.49 (s, 9H).

C. 3-(4-Chloro-3-methyl-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

- 10 To a solution of 3-(4-chloro-3-methyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (18.26 g, 53 mmol) and epichlorohydrin (41.12 mL, 526 mmol) in DMF (100 mL) was added cesium carbonate (20.56 g, 63 mmol). The reaction mixture was allowed to stir for 72 h, diluted with EtOAc (200 mL) and washed with saturated NaHCO₃ and brine. The organic layer was dried over Na₂SO₄, concentrated and purified by column chromatography (silica, 20% acetone/CH₂Cl₂) to afford 3-(4-chloro-3-methyl-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (12.0 g, 57%). TLC (silica, 20% acetone/CH₂Cl₂): R_f = 0.68. MS (electrospray) *m/z* 491.2 (491.2, calculated for C₂₇H₃₁ClN₆O, [M+H]⁺). ¹H NMR (400 MHz, CDCl₃) 7.55 (s, 1H), 7.36 (m, 2H), 4.61 (m, 2H), 4.38-4.47 (m, 1H), 4.11 (dd, *J* = 14.3, 5.7 Hz, 1H), 3.67-3.79 (m, 2H), 3.34 (m, 1H), 2.83 (t, *J* = 4.5 Hz, 1H), 2.75 (m, 2H), 2.51 (m, 1H), 2.41 (s, 3H), 1.48 (s, 9H).

D. 3-(4-Chloro-3-methyl-phenyl)-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

- 30 3-(4-Chloro-3-methyl-phenyl)-1-oxiranylmethyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (5.33 g, 13.2 mmol) and 1-(2-cyanophenyl)-piperazine (2.97 g, 15.86 mmol) were partially dissolved in EtOH (50 mL) and triethylamine (2 mL). The reaction mixture was heated to 80 °C for 18 h. The mixture was concentrated and purified by column

chromatography (silica, 20% acetone/CH₂Cl₂) to give 3-(4-chloro-3-methyl-phenyl)-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (6.51 g, 83%) as a yellow solid. TLC (silica, 20% acetone/CH₂Cl₂): *R_f* = 0.35. MS (electrospray): *m/z* 591.3 (591.3, calculated for C₃₂H₃₉ClN₆O₃, [M+H]⁺).

E. 2-(4-{3-[3-(4-Chloro-3-methyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile.

3-(4-Chloro-3-methyl-phenyl)-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (1.26 g, 2.13 mmol) was dissolved in trifluoroacetic acid (3 mL) and CH₂Cl₂ (3 mL) and allowed to stir for 2 h. The reaction mixture was concentrated, taken up in EtOAc (50 mL) and washed with aqueous NaHCO₃ (2 X 25 mL). The EtOAc layer was dried over Na₂SO₄ and concentrated to give 2-(4-{3-[3-(4-chloro-3-methyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile (1.05 g, 99%) as a yellow foam. TLC (silica, 10% MeOH/CH₂Cl₂): *R_f* = 0.27. MS (electrospray): *m/z* 491.2 (491.2, calculated for C₂₇H₃₁ClN₆O, [M+H]⁺). ¹H NMR (400 MHz, CDCl₃): 9.8 (br s, 1H), 7.55 (d, *J* = 7.6 Hz, 1H), 7.50 (t, *J* = 8.2 Hz, 1H), 7.38 (s, 1H), 7.31 (d, *J* = 8.2 Hz, 1H), 7.20 (d, *J* = 8.2 Hz, 1H), 7.11 (t, *J* = 8.2 Hz, 1H), 6.98 (d, *J* = 8.2 Hz, 1H), 4.56 (br s, 1H), 4.12-4.32 (m, 4H), 2.98-3.51 (m, 13 H), 2.35 (s, 3H).

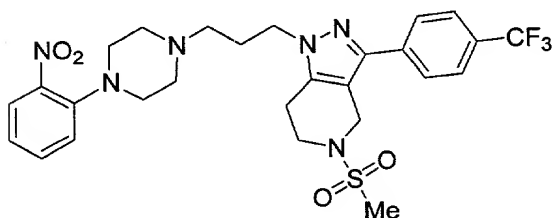
F. (3-(4-Chloro-3-methyl-phenyl)-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-oxo-acetic acid methyl ester.

2-(4-{3-[3-(4-Chloro-3-methyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile (58 mg, 0.118 mmol) was dissolved in CH₂Cl₂ (0.59 mL) and treated with methyl chlorooxoacetate (16 mg, 0.129 mmol). The reaction mixture was allowed to stir for 18 h at room temperature. Column chromatography (silica, 2-10% MeOH/CH₂Cl₂) gave (3-(4-chloro-3-methyl-phenyl)-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-oxo-acetic acid methyl ester (54 mg, 79%) as a white solid. MS (electrospray): *m/z* 577.3 (577.2,

calculated for $C_{30}H_{33}ClN_6O_4$, $[M+H]^+$. 1H NMR (400 MHz, $CDCl_3$): 7.32-7.62 (m, 5H), 7.14 (t, $J = 7.6$ Hz, 1H), 7.05 (d, $J = 8.2$ Hz, 1H), 4.59-4.80 (m, 3H), 4.12-4.28 (m, 2H), 3.92 (s, 3H), 3.78-3.86 (m, 2H), 3.44-3.60 (m, 5H), 3.15-3.40 (m, 4H), 2.83-3.05 (m, 2H), 2.41 (s, 3H).

5

EXAMPLE 16



10 5-Methanesulfonyl-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine.

A. 1-Methanesulfonyl-piperidin-4-one.

15 Potassium carbonate (324 g, 2340 mmol) was added to a solution of 4-piperidone monohydrate hydrochloride (90 g, 586 mmol) in chloroform (300 mL) and water (300 mL). The slurry was cooled to 0 °C and treated with methanesulfonyl chloride (136 mL, 1760 mmol) by dropwise addition over a 1 h period (gas evolution was observed). The reaction mixture was allowed to

20 shake for 72 h and was partitioned between CH_2Cl_2 (500 mL) and saturated aqueous $NaHCO_3$ (500 mL). The aqueous layer was extracted with CH_2Cl_2 (3 X 200 mL). The organic layer was washed with 1% $KHSO_4$ (250 mL), dried (Na_2SO_4), and concentrated to afford 90.5 g (87%) of a white solid. MS (electrospray): exact mass calculated for $C_6H_{11}NO_3S$, 177.1; m/z found, 178.1

25 $[M+H]^+$. HPLC (reverse phase conditions): $t_R = 2.19$ min. 1H NMR (400 MHz, $CDCl_3$): 3.60 (t, $J = 6.5$ Hz, 4H), 2.89 (s, 3H), 2.59 (t, $J = 6.3$ Hz, 4H).

B. 5-Methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine.

p-Toluenesulfonic acid (1.34 g, 7.0 mmol) and morpholine (25.83 mL, 296 mmol) were added to a solution of 1-methanesulfonyl-piperidin-4-one (50.0 g, 282 mmol) in benzene (282 mL). The reaction mixture was heated in a flask equipped with a condenser and a Dean-Stark trap at reflux for 15 h. The reaction mixture was cooled and concentrated in vacuo to give the enamine which was used without further purification. The enamine was dissolved in CH₂Cl₂ (200 mL) and cooled to 0 °C. To this was added triethylamine (47.2 mL, 339 mmol) followed by dropwise addition of 4-trifluoromethylbenzoyl chloride (42.3 mL, 285 mmol) dissolved in CH₂Cl₂ (82 mL). The reaction mixture was allowed to warm to room temperature and stirred for 20 h. The reaction mixture was washed with 1 *N* aqueous HCl (250 mL) and the CH₂Cl₂ layer was separated, dried (Na₂SO₄), and concentrated. The resulting oil was taken up in EtOH (300 mL) and treated with hydrazine (44.3 mL, 1.41 mol) at 0 °C. The reaction mixture was allowed to warm to room temperature and stirred for 24 h. The mixture was concentrated and the resulting solid was filtered with EtOH wash and dried in vacuo to afford 70 g (72%) of 5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-*c*]pyridine as a white solid. MS (electrospray): exact mass calculated for C₁₄H₁₄F₃N₃O₂S, 345.0; *m/z* found, 346.0 [M+H]⁺. HPLC (reverse phase conditions): *t*_R = 6.33 min. ¹H NMR (400 MHz, CDCl₃): 7.72 (s, 4H), 4.58 (s, 2H), 3.69 (t, *J* = 5.7 Hz, 2H), 2.99 (t, *J* = 5.7 Hz, 2H), 2.92 (s, 3H).

C. 3-[5-Methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-*c*]pyridin-1-yl]-propan-1-ol.

Cs₂CO₃ (33.74 g, 103.5 mmol) was added to a solution of 5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-*c*]pyridine (29.8 g, 86.3 mmol) in anhydrous DMF (70 mL) and stirred for 25 min. 3-Bromo-1-propanol (8.6 mL, 13.2 g, 94.9 mmol) was added and stirred under N₂ at room temperature for 18 h. Water (500 mL) was added to the reaction and stirred for 5 min. The precipitated material was filtered out and washed with water (4 X 100 mL) and dried in a Freeze Drying System. The crude material (31.0 g) was taken up in anhydrous DMF (65 mL) and Cs₂CO₃ (33.74 g, 103.5 mmol) was added, and stirred for 10 min. 3-Bromo-1-propanol (8.6 mL, 13.2 g, 94.9 mmol)

and MeOH (6.0 mL, 4.75 g, 148 mmol) were added and stirring continued under N₂ at room temperature for 15 h. Water (500 mL) was added to the reaction and stirred for 10 min. The precipitated material was filtered and washed with water (3 X 100 mL). The filter cake was dissolved in CH₂Cl₂ (200 mL) and washed with brine (50 mL), dried (Na₂SO₄), and concentrated. The solid was triturated with Et₂O (200 mL), filtered, washed with Et₂O, and dried to furnish 16.0 g of the desired compound. The mother liquor was chromatographed (silica, 0-10% acetone/EtOAc) to obtain an additional 3.0 g of the title compound. The combined yield was 54.6%. MS (electrospray): calculated for C₁₇H₂₀F₃N₃O₃S, 403.12; *m/z* found, 404.0 [M+H]⁺, 426.0 [M+Na]⁺. ¹H NMR (400 MHz, CDCl₃): 7.71 (d, *J* = 8.2 Hz, 2H), 7.66 (d, *J* = 8.5 Hz, 2H), 4.55 (s, 2H), 4.23 (t, *J* = 6.5 Hz, 2H), 3.70-3.63 (m, 4H), 2.90 (s, 3H), 2.90 (t, *J* = 5.1 Hz, 2H), 2.62 (t, *J* = 5.9 Hz, 1H), 2.06 (q, *J* = 6.1 Hz, 2H).

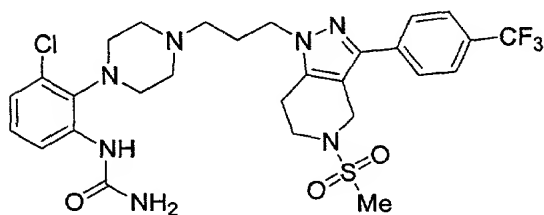
D. 3-[5-Methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propionaldehyde.

Dess-Martin periodinane (3.45 g, 8.2 mmol) was added to a solution of 3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-1-ol (3.0 g, 7.4 mmol) in CH₂Cl₂ (20 mL) at 0 °C under N₂. After 15 min, the reaction was allowed to warm to room temperature and stirred for another 1.5 h. The reaction was diluted with Et₂O (60 mL) and 20 % aq. NaHCO₃ (35 mL) was added slowly. Then Na₂S₂O₃ was added and stirred at room temperature for 30 min. The layers were separated and the aqueous portion was extracted with Et₂O (2 x 30 mL). The combined organic extracts were washed with brine, dried (Na₂SO₄) and concentrated. MPLC (1-10% MeOH/CH₂Cl₂) afforded 2.53 g of the desired aldehyde in 85% yield. MS (electrospray): calculated for C₁₇H₁₈F₃N₃O₃S, 401.11; *m/z* found, 402.1 [M+H]⁺, 434.1 [M+MeOH+H]⁺. ¹H NMR (400 MHz, CDCl₃): 9.82 (s, 1H), 7.63 (d, *J* = 8.4 Hz, 2H), 7.58 (d, *J* = 8.4 Hz, 2H), 4.68 (s, 2H), 4.25 (t, *J* = 6.1 Hz, 2H), 3.63 (t, *J* = 5.8 Hz, 4H), 3.14 (t, *J* = 6.1 Hz, 2H), 2.92 (t, *J* = 5.8 Hz, 2H), 2.81 (s, 3H).

E. 5-Methanesulfonyl-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine.

To a stirred solution of 3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propionaldehyde (0.060 g, 0.15 mmol) and 1-(2-nitro-phenyl)-piperazine (0.032 g, 0.157 mmol) in CH₂Cl₂ (0.5 mL), glacial AcOH (8.5 μ L, 0.15 mmol) was added and stirred for 15 min at room temperature. NaBH(OAc)₃ (0.041 g, 0.19 mmol) was added and stirred under nitrogen overnight. Saturated NaHCO₃ (0.5 mL) was then added and stirred for 15 min. The layers separated and the aqueous layer was extracted with CH₂Cl₂ (0.5 mL). MPLC purification (silica, 2-15% MeOH/CH₂Cl₂) afforded the desired product as a white solid (0.063 g, 71%). TLC (silica, 12% MeOH/CH₂Cl₂): R_f = 0.67. MS (electrospray): exact mass calculated for C₂₇H₃₁F₃N₆O₄S, 592.21; *m/z* found, 593.2 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃): 7.80 (dd, *J* = 1.6, 8.2 Hz, 1H), 7.77 (d, *J* = 8.3 Hz, 2H), 7.70 (d, *J* = 8.3 Hz, 2H), 7.52 (ddd, *J* = 1.6, 7.3, 8.3 Hz, 1H), 7.19 (dd, *J* = 1.2, 8.3 Hz, 1H), 7.09 (m, 1H), 4.59 (s, 2H), 4.17 (t, *J* = 6.9 Hz, 2H), 3.71 (t, *J* = 5.8 Hz, 2H), 3.13 (br t, *J* = 4.8 Hz, 4H), 2.96 (t, *J* = 5.6 Hz, 2H), 2.95 (s, 3H), 2.66 (br t, *J* = 4.4 Hz, 4H), 2.51 (t, *J* = 7.0 Hz, 2H), 2.17 (q, *J* = 6.9 Hz, 2H).

EXAMPLE 17



1-[3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-3-methyl-urea.

A. 4-(2-chloro-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester.

To a stirred solution of 1,2-dichloro-3-nitrobenzene (0.96 g, 5.0 mmol) and piperazine-1-carboxylic acid tert-butyl ester (0.93 g, 5.0 mmol) in acetonitrile (5 mL) was added of K_2CO_3 (1.38 g, 10 mmol). The mixture was heated at reflux for 48 h. The solvent was removed under reduced pressure. The crude material was partitioned between EtOAc (100 mL) and H_2O (20 mL). The organic layer was washed with H_2O (2 x 20 mL), dried over Na_2SO_4 and concentrated. Column chromatography (silica, 10-20% EtOAc/hexanes) provided 4-(2-chloro-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester (1.2 g, 70%). TLC (silica, 20% EtOAc/hexanes): R_f = 0.45. MS (electrospray): exact mass calculated for $C_{15}H_{20}ClN_3O_4$, 341.1; m/z found, 364.1 $[M+Na]^+$. 1H NMR (400 MHz, $CDCl_3$) 7.56 (dd, J = 8.2, 1.4 Hz, 1H), 7.50 (dd, J = 8.2, 1.4 Hz, 1H), 7.13 (t, J = 8.2 Hz, 1H), 3.38-3.56 (m, 4H), 3.06 (m, 4H), 1.48 (s, 9H).

B. 1-(2-chloro-6-nitro-phenyl)-piperazine.

4-(2-Chloro-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester (1.87 g, 5.47 mmol) was dissolved in trifluoroacetic acid (5.0 mL) and CH_2Cl_2 (5.0 mL) and allowed to stir for 2 h. The reaction mixture was concentrated, diluted with EtOAc, and washed with saturated aq. $NaHCO_3$. The organic layer was dried over Na_2SO_4 , concentrated and purified by column chromatography (silica, 100% CH_2Cl_2) to afford 1-(2-chloro-6-nitro-phenyl)-piperazine (1.26 g, 95%). MS (electrospray): exact mass calculated for $C_{10}H_{12}ClN_3O_2$, 241.1; m/z found, 242.1 $[M+H]^+$. 1H NMR (400 MHz, $CDCl_3$): 7.54 (dd, J = 8.2, 1.6 Hz, 1H), 7.49 (dd, J = 8.2, 1.6 Hz, 1H), 7.10 (t, J = 8.2 Hz, 1H), 3.08 (br s, 4H), 2.99 (br s, 4H), 2.07-2.12 (m, 1H).

C. 1-{3-[4-(2-Chloro-6-nitro-phenyl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine.

To a stirred solution of 3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propionaldehyde (0.5 g, 1.25 mmol) and 1-(2-chloro-6-nitrophenyl)-piperazine (0.301 g, 1.25 mmol) in CH_2Cl_2 (6 mL) was added sodium sulfate (0.354 g, 2.50 mmol) and sodium triacetoxymethylborohydride (0.396 g, 1.87 mmol). The mixture was allowed to stir at

room temperature overnight. The mixture was diluted with CH_2Cl_2 and washed with water. The CH_2Cl_2 layer was dried over Na_2SO_4 , and the solvent was removed under reduced pressure. The residue was purified by column chromatography (silica, 10% acetone/ CH_2Cl_2) to afford of 1-{3-[4-(2-chloro-6-nitro-phenyl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine (0.380 g, 49%). MS (electrospray): exact mass calculated for $\text{C}_{27}\text{H}_{30}\text{ClF}_3\text{N}_6\text{O}_4\text{S}$, 626.2; m/z found, 627.2 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3): 7.73 (d, $J = 8.2$ Hz, 2H), 7.66 (d, $J = 8.2$ Hz, 2H), 7.54 (dd, $J = 8.2, 1.2$ Hz, 1H), 7.49 (dd, $J = 8.2, 1.2$ Hz, 1H), 7.10 (t, $J = 8.2$ Hz, 1H), 4.58 (s, 2H), 4.13 (t, $J = 6.5$ Hz, 2H), 3.71 (t, $J = 5.9$ Hz, 2H), 3.01-3.11 (m, 4H), 2.95 (t, $J = 5.9$ Hz, 2H), 2.92 (s, 3H), 2.42-2.53 (m, 4H), 2.40 (t, $J = 6.5$ Hz, 2H), 2.12 (q, $J = 6.5$ Hz, 2H).

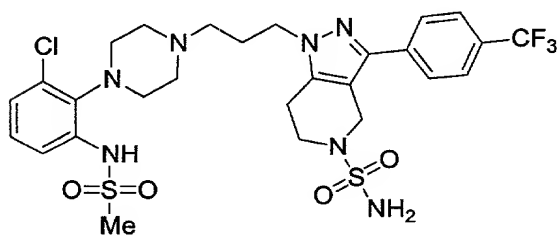
D. 3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenylamine.

To a stirred solution of 1-{3-[4-(2-chloro-6-nitro-phenyl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine (0.153 g, 0.244 mmol) in EtOH (2.44 mL) was added zinc dust (0.80 mg, 1.22 mmol) and slow addition of acetic acid (0.70 mL). After 15 min the yellow solution became colorless and the excess zinc dust was filtered through a plug of celite. The filtrate was concentrated and the residue was purified by column chromatography (silica, 0-10% MeOH/ CH_2Cl_2) to afford 3-chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenylamine (0.146 g, 100%). MS (electrospray): exact mass calculated for $\text{C}_{27}\text{H}_{32}\text{ClF}_3\text{N}_6\text{O}_2\text{S}$, 596.2; m/z found, 597.2 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3): 7.73 (d, $J = 8.2$ Hz, 2H), 7.66 (d, $J = 8.2$ Hz, 2H), 6.88 (t, $J = 8.2$ Hz, 1H), 6.63 (t, $J = 7.6$ Hz, 2H), 4.55 (s, 2H), 4.36 (s, 2H), 4.15 (t, $J = 6.5$ Hz, 2H), 3.60-3.70 (m, 4H), 2.97 (t, $J = 5.3$ Hz, 2H), 2.90 (s, 3H), 2.83 (d, $J = 10.8$ Hz, 2H), 2.74 (d, $J = 11.5$ Hz, 2H), 2.37 (t, $J = 6.6$ Hz, 2H), 2.11-2.20 (m, 4H).

E. 1-[3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-3-methyl-urea.

To a stirred solution of 3-chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenylamine (0.062 g, 0.104 mmol) in CH_2Cl_2 (0.52 mL) was added trimethylsilyl isocyanate (0.017 mL, 0.125 mmol). The reaction mixture was allowed to stir for 48 h at room temperature. The reaction had not gone to completion, so an additional 0.017 mL (0.125 mmol) of trimethylsilyl isocyanate was added and the reaction was heated to 45 °C for 10 h. Column chromatography (silica, 3-10% MeOH/ CH_2Cl_2) afforded 1-[3-chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-3-methyl-urea (0.015 g, 22%). MS (electrospray): exact mass calculated for $\text{C}_{28}\text{H}_{33}\text{ClF}_3\text{N}_7\text{O}_3\text{S}$, 639.2; m/z found, 640.2 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3): 8.26 (br s, 1H), 8.05 (d, J = 8.2 Hz, 1H), 7.73 (d, J = 8.2 Hz, 2H), 7.66 (d, J = 8.2 Hz, 2H), 7.09 (t, J = 8.2 Hz, 1H), 6.92 (d, J = 8.2 Hz, 1H), 4.65 (s, 2H), 4.55 (s, 2H), 4.15 (t, J = 6.7 Hz, 2H), 3.65-3.73 (m, 4H), 2.96 (t, J = 5.6 Hz, 2H), 2.87-2.92 (m, 2H), 2.91 (s, 3H), 2.70 (d, J = 11.4 Hz, 2H), 2.40 (t, J = 6.7 Hz, 2H), 2.09-2.22 (m, 4H).

EXAMPLE 18



1-{3-[4-(2-Chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-sulfonic acid amide.

A. 3-(4-Trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

To a stirred solution of 500 g (2.51 mol) of 1-*tert*-butoxycarbonyl-4-piperidone and 87.1 g (2.76 mol) of morpholine in benzene (1.25 L) was added a catalytic amount (~ 0.25 g) of *p*-TsOH. The mixture was heated to reflux for 36 h with a Dean-Stark trap. One half of the solvent was removed under reduced pressure and the resulting solution was cooled and filtered. The filtrate was then concentrated to yield 630 g (94%) of an orange red oil. The enamine was divided and 320 g (1.19 mol) was diluted with CH₂Cl₂ (1.0 L) and 165.0 mL (1.19 mol) of Et₃N was added. The mixture was cooled to 0 °C and a solution of 225 g (1.08 mol) of 4-trifluoromethylbenzoyl chloride in CH₂Cl₂ (0.5 L) was added slowly by dropping funnel over 1h. The mixture was allowed to warm to rt and stir overnight. The reaction was then diluted with 1 *N* HCl (450 mL) and stirred vigorously for 3 h. The aqueous layer was extracted with CH₂Cl₂ (3 x 500 mL) and the combined extracts were dried over Na₂SO₄ and the solvent was removed under reduced pressure. The crude oil was diluted with EtOH (1 L) and cooled to 0 °C. To this stirred solution was slowly added 115 g (3.57 mol) of hydrazine and the mixture was allowed to warm to rt and stir overnight during which time a white precipitate formed. The volume of the reaction was reduced to ~500 mL and cooled. The precipitate was collected to afford 285 g (72% from enamine) of a white solid. ¹H NMR (400 MHz, CDCl₃): 7.63-7.55 (m, 4H), 4.58 (br s, 2H), 3.69-3.62 (br m, 2H), 2.74-2.68 (br m, 2H), 1.47 (s, 9H).

B. 1-(2-Methoxycarbonyl-ethyl)-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

3-(4-Trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (1.85g, 5.04 mmol) and methyl acrylate (0.50 mL, 5.6 mmol) were combined in toluene (30 mL) and heated to 75 °C. The resulting mixture was treated with *t*-BuONa (100 mg), and heating continued for 48 h. The mixture was allowed to cool and partitioned between EtOAc (300 mL) and NaHCO₃ (75 mL). The aqueous layer was extracted with EtOAc (3 x 75 mL). The combined extracts were dried over Na₂SO₄ and concentrated.

Column chromatography (silica, 30-60% EtOAc/hexanes) afforded 343 mg (15%) of the title compound. TLC (silica, 50% EtOAc/hexanes): R_f = 0.4. MS (electrospray): m/z calculated for $C_{22}H_{27}F_3N_3O_4$ [$M^+ + H$] 454.20, found 454.1. 1H NMR ($CDCl_3$, 400 MHz): 7.75 (br d, J = 8.1 Hz, 2H), 7.64 (br s, 2H), 4.63 (br s, 2H), 4.30 (t, J = 6.6 Hz, 2H), 3.75 (br s, 2H), 3.68 (s, 3H), 2.98 (t, J = 6.6 Hz, 2H), 2.79 (br t, J = 5.6 Hz, 2H), 1.48 (s, 9H).

C. 1-(3-Hydroxy-propyl)-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester .

A solution of $LiBH_4$ (26 mg, 1.2 mmol) in THF (0.5 mL) was added to a 0 °C solution of 1-(2-methoxycarbonyl-ethyl)-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (317 mg, 0.70 mmol) in THF (4.0 mL). The mixture was stirred for 5 min then additional $LiBH_4$ (15 mg) was added and stirring continued for 17 h. The mixture was partitioned between EtOAc (80 mL) and saturated aqueous $NaHCO_3$ (20 mL). The aqueous layer was extracted with EtOAc (2 x 20 mL). The combined extracts were dried over Na_2SO_4 and concentrated. Column chromatography (silica, 0-8% MeOH/ CH_2Cl_2) afforded 268 mg (95%) of the title compound. HPLC (reverse phase conditions), t_R = 6.82 min. MS (electrospray): m/z calculated for $C_{21}H_{26}F_3N_3O_3$ [$M^+ + Na$] 448.18, found 448.10. 1H NMR ($CDCl_3$, 400 MHz): 7.73 (br d, J = 8.2 Hz, 2H), 7.65 (br s, 2H), 4.64 (br s, 2H), 4.21 (t, J = 6.4 Hz, 2H), 3.76 (br s, 2H), 3.66 (t, J = 5.7 Hz, 2H), 2.73 (br t, J = 5.4 Hz, 2H), 2.04 (q, J = 6.1, 2H), 1.48 (s, 9H).

D. 1-(3-Oxo-propyl)-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

Dess-Martin periodinane (1.43 g, 3.36 mmol) was added portion wise to a stirred solution of 1-(3-hydroxy-propyl)-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydropyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (1.30 g, 3.05 mmol) in CH_2Cl_2 (15 mL) at 0 °C under N_2 . Then the reaction was stirred at 0 °C for 15 min and allowed to warm to room temperature. After stirring at room temperature for 1.5 h the reaction was diluted with Et_2O (50 mL) and saturated $NaHCO_3$ (15 mL) was added slowly (caution! gas evolution). Then

Na₂S₂O₃·5H₂O (5.31 g, 21.4 mmol) was added and stirred for 30 min. The layers were separated and the aqueous layer was extracted with Et₂O (2 x 30 mL). The combined extracts were washed with brine, dried (Na₂SO₄) and concentrated. MPLC (1-10% MeOH/CH₂Cl₂) afforded the aldehyde in 79% yield (1.02 g). TLC (silica, 10% MeOH/CH₂Cl₂): R_f = 0.67. MS (electrospray) calculated for C₂₁H₂₄F₃N₃O₃, 424.2 ([M+H]⁺), *m/z* found, 424.2. ¹H NMR (400 MHz, CDCl₃): 9.82 (s, 1H), 7.65 (br d, *J* = 8.0 Hz, 2H), 7.54 (br s, 2H), 4.53 (s, 2H), 4.21 (t, *J* = 6.2 Hz, 2H), 3.68 (br s, 2H), 3.04 (t, *J* = 6.2 Hz, 2H), 2.70 (t, *J* = 5.6 Hz, 2H), 1.39 (s, 9H).

E. 4-(2-Chloro-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester.

To a stirred solution of 0.96 g (5.0 mmol) of 1,2-dichloro-3-nitrobenzene and 0.93 g (5.0 mmol, 1 eq) of 1-*tert*-butoxycarbonylpiperazine in acetonitrile (5 mL) was added 1.38 g (10 mmol, 2 eq) of K₂CO₃. The mixture was heated to reflux for 48 h. The solvent was removed under reduced pressure. The crude product was partitioned between EtOAc (100 mL) and 20 mL of H₂O. The organic layer was washed with H₂O (2 x 20 mL), dried over Na₂SO₄ and concentrated. Column chromatography (silica, 10-20% EtOAc/hexanes) provided 1.2 g (70%) of 4-(2-chloro-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester. TLC (silica, 20% EtOAc/hexanes): R_f = 0.45. ¹H NMR (400 MHz, CDCl₃): 7.56 (dd, *J* = 8.2, 1.4 Hz, 1H), 7.50 (dd, *J* = 8.2, 1.4 Hz, 1H), 7.13 (t, *J* = 8.2 Hz, 1H), 3.56-3.38 (m, 4H), 3.10-3.00 (m, 4H), 1.48 (s, 9H).

F. 1-{3-[4-(2-Chloro-6-nitro-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

4-(2-Chloro-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester (940 mg, 2.75 mmol) in 10 mL of CH₂Cl₂ was treated with 5 mL of trifluoroacetic acid and stirred at 25 °C for 1 h. The volatiles were then removed. The residue was taken up in CH₂Cl₂ (60 mL) and KOH (4 N, 20 mL). The organic layer was separated, dried over Na₂SO₄, and concentrated. The yellow oil was dissolved in CH₂Cl₂ and added into the 996 mg (2.35 mmol) of 1-(3-oxo-propyl)-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxy

lic acid tert-butyl ester. The yellow solution was treated with glacial acetic acid (0.8 mL, 6 eq) and stirred at 25 °C for 1 h. NaBH(OAc)₃ (1.5 g, 7.05 mmol) was added and stirred under nitrogen for 2 h. Then saturated NaHCO₃ (20 mL) was added and stirred for 30 min, and the layers were separated. The organic
 5 extract was washed with brine, dried over Na₂SO₄, and concentrated under reduced pressure. Column chromatography (silica, 2-5% MeOH/CH₂Cl₂) afforded 1-{3-[4-(2-chloro-6-nitro-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester as a white solid (1.40 g, 92%). TLC (silica, 5%
 10 MeOH/CH₂Cl₂): R_f = 0.3. MS (electrospray): exact mass calculated for C₃₁H₃₆ClF₃N₆O₄, 648.24; *m/z* found 649.3 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃): 7.69 (d, *J* = 8.2 Hz, 1H), 7.60-7.50 (m, 1H), 7.45-7.37 (m, 4H), 7.02 (t, *J* = 8.2 Hz, 1H), 4.58 (br s, 2H), 4.04 (t, *J* = 6.7 Hz, 2H), 3.73-3.65 (m, 2H), 3.05-2.95 (m, 4H), 2.71 (t, *J* = 5.6 Hz, 2H), 2.50-2.35 (m, 4H), 2.30 (t, *J* = 6.8 Hz, 2H),
 15 2.05-1.95 (m, 2H), 1.41 (s, 9H).

G. 1-{3-[4-(2-Amino-6-chloro-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

20 A solution of 360 mg (0.56 mmol) of 1-{3-[4-(2-chloro-6-nitro-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester in 4 mL of MeOH was treated with 182 mg (5 eq) of zinc dust and glacial acetic acid (1.57 mL, 50 eq) at 25 °C. The reaction mixture was stirred at 25 °C for 1 h. The reaction mixture was
 25 then filtered through a pad of celite and concentrated to obtain a thick oil. The residue was taken up in CH₂Cl₂ (50 mL) and sat. NaHCO₃ (20 mL). The organic layer was separated, washed with H₂O (2x 10 mL), dried over Na₂SO₄, and concentrated to afford 1-{3-[4-(2-amino-6-chloro-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-
 30 carboxylic acid tert-butyl ester. TLC (silica, 10% MeOH/CH₂Cl₂): R_f = 0.3. MS (electrospray): exact mass calculated for C₃₁H₃₈ClF₃N₆O₂, 618.27; *m/z* found, 619.3 [M+H]⁺.

H. 1-{3-[4-(2-Chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester.

A solution of 1-{3-[4-(2-amino-6-chloro-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester (257 mg, 0.42 mmol) in 4 mL of CH₂Cl₂ was treated with 32 L (0.42 mmol, 1.0 eq) of methanesulfonyl chloride and 116 L (0.83 mmol, 2 eq) of triethylamine and the reaction mixture stirred at 25 °C for 1 h. EtOAc (40 mL) and sat. NaHCO₃ (20 mL) were added. The organic layer was separated and washed with H₂O (20 mL), brine (20 mL), dried over Na₂SO₄, and concentrated to afford the crude 1-{3-[4-(2-chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester. TLC (silica, 10% MeOH/CH₂Cl₂): R_f = 0.3. MS (electrospray): exact mass calculated for C₃₂H₄₀ClF₃N₆O₄S, 696.25; *m/z* found, 697.2 [M+H]⁺.

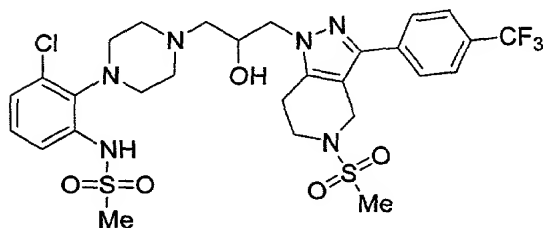
I. 1-{3-[4-(2-Chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-tert-butoxycarbonyl-sulfonic acid amide.

A solution of 97 mg (0.14 mmol) of 1-{3-[4-(2-chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester in 3 mL of CH₂Cl₂ was treated with 1.5 mL of trifluoroacetic acid. The reaction mixture was stirred at 25 °C for 1 h before all volatiles were removed. To this crude material in 0.5 mL of CH₂Cl₂ was added dropwise a premixed solution of chlorosulfonyl isocyanate (18 μL, 0.209 mmol) and 2-methyl-2-propanol (20 μL, 0.209 mmol) in CH₂Cl₂ (0.150 mL). The reaction mixture was allowed to stir at 25 °C overnight. Preparative TLC (silica, 2-10% MeOH/CH₂Cl₂) provided the title compound (84 mg, 78%). TLC (silica, 10% MeOH/CH₂Cl₂): R_f = 0.3. MS (electrospray): exact mass calculated for C₃₂H₄₁ClF₃N₇O₆S₂, 775.22; *m/z* found, 776.2 [M+H]⁺.

J. 1-{3-[4-(2-Chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-sulfonic acid amide.

1-{3-[4-(2-Chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-tert-butoxycarbonyl-sulfonic acid amide (84 mg, 0.11 mmol) was dissolved in trifluoroacetic acid (0.75 mL) and CH₂Cl₂ (0.75 mL). The reaction mixture was allowed to stir at 25 °C for 2 h. Removal of volatiles under a stream of nitrogen provided 1-{3-[4-(2-chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-sulfonic acid amide in quantitative yield as a trifluoroacetic acid salt. MS (electrospray): exact mass calculated for C₂₇H₃₃ClF₃N₇O₄S₂, 675.17; *m/z* found, 676.2 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃): 7.73 and 7.63 (AB pattern, *J* = 8.2 Hz, 4H), 7.37 (d, *J* = 7.8 Hz, 1H), 7.13 (t, *J* = 7.8 Hz, 1H), 7.04 (d, *J* = 7.8 Hz, 1H), 4.32 (s, 2H), 4.20 (t, *J* = 6.3 Hz, 2H), 3.87-3.80 (m, 2H), 3.80-3.75 (m, 4H), 3.70-3.25 (m, 7H), 3.00-2.75 (m, 4H), 2.25-2.15 (m, 2H).

EXAMPLE 19



N-[3-Chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-methanesulfonamide.

A. 5-Methanesulfonyl-1-oxiranylmethyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine.

5-Methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine (10.0 g, 29.0 mmol) and epichlorohydrin (24 mL, 307

mmol) were set stirring in DMF (150 mL) containing Cs_2CO_3 (10.4 g, 31.9 mmol). After stirring at room temperature for 4 days the mixture was evaporated, brought up in EtOAc and washed with water. The organics were dried (MgSO_4) and evaporated to give a light yellow solid. Column chromatography (silica, 5% acetone/ CH_2Cl_2) gave 4.1g (35%) of a white solid. TLC (silica, 5% acetone/ CH_2Cl_2): R_f = 0.28. MS (electrospray): exact mass calculated for $\text{C}_{17}\text{H}_{18}\text{F}_3\text{N}_3\text{O}_3\text{S}$, 401.10; m/z found, 402.1 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3): 7.84 (d, J = 8.3 Hz, 2H), 7.79 (d, J = 8.3 Hz, 2H), 4.70-4.62 (m, 3H), 4.25 (d, J = 5.4 Hz, 1H), 3.90-3.70 (m, 2H), 3.47 (m, 1H), 3.10-2.9 (m, 6H), 2.65-2.60 (m, 1H).

B. 4-(2-Chloro-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester.

To a stirred solution of 0.96 g (5.0 mmol) of 1,2-dichloro-3-nitrobenzene and 0.93 g (5.0 mmol, 1 eq) of 1-*tert*-butoxycarbonylpiperazine in acetonitrile (5 mL) was added 1.38 g (10 mmol, 2 eq) of K_2CO_3 . The mixture was heated to reflux for 48 h. The solvent was removed under reduced pressure. The crude product was partitioned between EtOAc (100 mL) and 20 mL of H_2O . The organic layer was washed with H_2O (2 x 20 mL), dried over Na_2SO_4 and concentrated. Column chromatography (silica, 10-20% EtOAc/hexanes) provided 1.2 g (70%) of 4-(2-chloro-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester. TLC (silica, 20% EtOAc/hexanes): R_f = 0.45. ^1H NMR (400 MHz, CDCl_3): 7.56 (dd, J = 8.2, 1.4 Hz, 1H), 7.50 (dd, J = 8.2, 1.4 Hz, 1H), 7.13 (t, J = 8.2 Hz, 1H), 3.56-3.38 (m, 4H), 3.10-3.00 (m, 4H), 1.48 (s, 9H).

C. 4-(2-Amino-6-chloro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester.

A solution of 342 mg (1 mmol) of 4-(2-chloro-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester in 5.0 mL of MeOH was treated with 630 mg (10 mmol, 10 eq) of ammonium formate and a catalytic amount of 10% Pd-C (34 mg). The reaction mixture was stirred at 65 °C for 30 min. The reaction mixture was then filtered through a pad of celite and concentrated to obtain a yellow solid. TLC (silica, 5% acetone/ CH_2Cl_2): R_f = 0.40. MS (electrospray): exact mass calculated for $\text{C}_{15}\text{H}_{22}\text{ClN}_3\text{O}_2$, 311.14; m/z found, 312.1 $[\text{M}+\text{H}]^+$.

D. 4-(2-Chloro-6-methanesulfonylamino-phenyl)-piperazine-1-carboxylic acid tert-butyl ester.

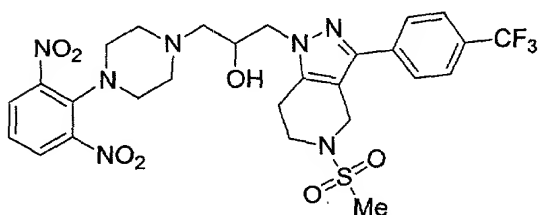
4-(2-Amino-6-chloro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester (163 mg, 0.53 mmol) in CH_2Cl_2 was treated with 62 μL (0.80 mmol, 1.5 eq) of methanesulfonyl chloride and 148 μL (1.06 mmol, 2 eq) of triethylamine and the reaction mixture stirred at 25 $^\circ\text{C}$ for 1 h. EtOAc (40 mL) and sat. NaHCO_3 (20 mL) were added. The organic layer was separated and washed with H_2O (20 mL), brine (20 mL), dried over Na_2SO_4 , and concentrated. Column chromatography (silica, 0-5% acetone/ CH_2Cl_2) provided 145 mg (70%) of 4-(2-chloro-6-methanesulfonylamino-phenyl)-piperazine-1-carboxylic acid tert-butyl ester. TLC (silica, 5% acetone/ CH_2Cl_2): R_f = 0.35. MS (electrospray): exact mass calculated for $\text{C}_{16}\text{H}_{24}\text{ClN}_3\text{O}_4\text{S}$, 389.12; m/z found, 388.1 (negative). ^1H NMR (400 MHz, CDCl_3): 7.41 (dd, J = 8.2, 1.6 Hz, 1H), 7.11 (t, J = 8.2 Hz, 1H), 6.99 (dd, J = 8.2, 1.6 Hz, 1H), 4.25-3.91 (m, 2H), 3.66-3.52 (m, 2H), 3.01 (s, 3H), 3.01-2.84 (m, 2H), 2.70-2.56 (m, 2H), 2.55-2.43 (m, 2H), 1.44 (s, 9H).

E. N-[3-Chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-methanesulfonamide.

4-(2-Chloro-6-methanesulfonylamino-phenyl)-piperazine-1-carboxylic acid tert-butyl ester (145 mg, 0.37 mmol) was dissolved in 3 mL of CH_2Cl_2 and treated with 1.5 mL of trifluoroacetic acid. The reaction mixture was stirred at 25 $^\circ\text{C}$ for 1 h before all volatiles were removed. The solid was treated with CH_2Cl_2 (20 mL) and aqueous KOH (4 N, 10 mL). The organic layer was separated, dried over Na_2SO_4 , and concentrated. The crude oil (90 mg) was dissolved in absolute EtOH (1.0 mL) and treated with 96 mg (0.24 mmol) of 5-methanesulfonyl-1-oxiranylmethyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine. The reaction mixture was refluxed at 85 $^\circ\text{C}$ for 3 h and then the solvent was removed. Column chromatography (silica, 0-5% MeOH/ CH_2Cl_2) provided 138 mg (20% over 4 steps) of N-[3-chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-methanesulfonamide. TLC (silica, 5% MeOH/ CH_2Cl_2): R_f = 0.45. MS

(electrospray): exact mass calculated for $C_{28}H_{34}ClF_3N_6O_5S_2$, 690.17; m/z found, 691.2 $[M+H]^+$. 1H NMR (400 MHz, $CDCl_3$): 8.28 (s, 1H), 7.65 and 7.59 (AB pattern, $J = 8.4$ Hz, 4H), 7.36 (d, $J = 8.1$ Hz, 1H), 7.07 (t, $J = 8.2$ Hz, 1H), 6.95 (d, $J = 8.2$ Hz, 1H), 4.54-4.44 (m, 2H), 4.21-3.94 (m, 3H), 3.77-3.52 (m, 4H),
 5 3.41 (m, 2H), 2.96 (s, 3H), 2.81 (s, 3H), 3.05-2.73 (m, 4H), 2.66-2.20 (m, 4H).

EXAMPLE 20



10 1-[4-(2,6-Dinitro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol.

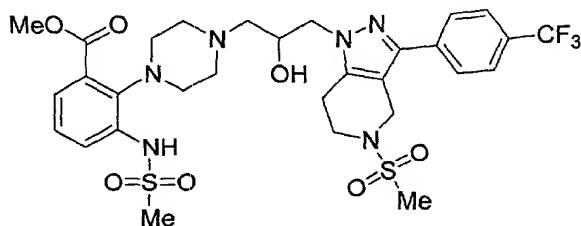
15 A. 4-(2,6-Dinitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester.

To a stirred solution of 1.01 g (5.0 mmol) of 1-chloro-2, 6-dinitrobenzene and 0.93 g (5.0 mmol) of 1-*tert*-butyloxycarbonylpiperazine in acetonitrile (5 mL) was added 1.38 g (10 mmol) of K_2CO_3 . The mixture was heated to reflux for 48 h. The solvent was removed under reduced pressure. The crude product was
 20 partitioned between EtOAc (100 mL) and 20 mL of H_2O . The organic layer was washed with H_2O (2 x 20 mL), dried over Na_2SO_4 and concentrated. Column chromatography (silica, 10-20% EtOAc/hexanes) provided 1.31 g (85%) of 4-(2,6-dinitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester
 TLC (silica, 20% EtOAc/hexanes): $R_f = 0.35$. 1H NMR (400 MHz, $CDCl_3$): 7.75
 25 (d, $J = 8.2$ Hz, 2H), 7.25 (t, $J = 8.2$ Hz, 1H), 3.30 (m, 4H), 2.95 (m, 2H), 1.44 (s, 9H).

B. 1-[4-(2,6-Dinitro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol.
 30 ol.

4-(2,6-Dinitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester (220 mg, 0.63 mmol) was dissolved in 5.0 mL of CH₂Cl₂ and treated with 3.0 mL of trifluoroacetic acid. The reaction mixture was stirred at 25 °C for 1 h before all volatiles were removed. The solid was treated with CH₂Cl₂ (20 mL) and aqueous KOH (4 N, 10 mL). The organic layer was separated, dried over Na₂SO₄, and concentrated. The crude oil (67 mg) was dissolved in absolute EtOH (1.2 mL) and treated with 141 mg (0.35 mmol, 1.3 eq) of 5-methanesulfonyl-1-oxiranylmethyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine. The reaction mixture was refluxed at 85 °C for 3 h and then the solvent was removed. Column chromatography purification (silica, 10-20% acetone/CH₂Cl₂) provided 150 mg (85%) of 1-[4-(2,6-dinitro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol. TLC (silica, 10% acetone/CH₂Cl₂): R_f = 0.3. MS (electrospray): exact mass calculated for C₂₇H₃₀F₃N₇O₇S, 653.19; *m/z* found, 654.2 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃): 7.71 (d, *J* = 8.2 Hz, 2H), 7.64 and 7.58 (AB pattern, *J* = 8.4 Hz, 4H), 7.20 (t, *J* = 8.2 Hz, 1H), 4.54 (s, 2H), 4.29-4.12 (m, 2H), 3.66 (t, *J* = 5.3 Hz, 2H), 3.70-2.95 (m, 9H), 2.91 (s, 3H), 2.67-2.32 (m, 4H).

20 EXAMPLE 21



25 2-(4-{2-Hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-3-methanesulfonylamino-benzoic acid methyl ester.

A. 4-(2-Methoxycarbonyl-6-nitro-phenyl)-piperazine-1-carboxylic acid tert-butyl ester.

To a stirred solution of 736 mg (2.83 mmol) of ethyl 2-bromo-3-nitrobenzoate and 579 mg (3.1 mmol, 1.1 eq) of 1-*tert*-butoxycarbonylpiperazine in *n*-butanol (6 mL) was added 330 mg (3.1 mmol, 1.1 eq) of Na₂CO₃. The mixture was heated to reflux for 4 h. The solvent was removed under reduced

- 5 pressure. The crude product was partitioned between EtOAc (100 mL) and 20 mL of H₂O. The organic layer was washed with H₂O (2 x 20 mL), dried over Na₂SO₄ and concentrated. Column chromatography (silica, 10-20% EtOAc/hexanes) provided 744 mg (72%) of 4-(2-methoxycarbonyl-6-nitro-phenyl)-piperazine-1-carboxylic acid *tert*-butyl ester. TLC (silica, 20% EtOAc/hexanes): R_f = 0.5. ¹H NMR (400 MHz, CDCl₃): 7.67 (dd, *J* = 8.2, 1.4 Hz, 1H), 7.62 (dd, *J* = 8.2, 1.4 Hz, 1H), 7.16 (t, *J* = 8.2 Hz, 1H), 3.86 (s, 3H), 3.44-3.36 (m, 4H), 3.03-2.95 (m, 4H), 1.48 (s, 9H).

15 B. 2-(4-{2-Hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4.3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-3-methanesulfonylamino-benzoic acid methyl ester.

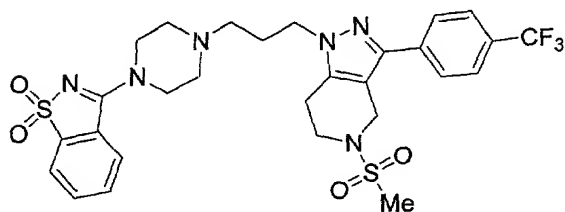
- A solution of 1.0 g (2.73 mmol) of 4-(2-methoxycarbonyl-6-nitro-phenyl)-piperazine-1-carboxylic acid *tert*-butyl ester in 18 mL of MeOH was treated with 893 mg (13.7 mmol, 5 eq) of zinc dust and glacial acetic acid (8 mL). The reaction mixture was stirred at 25 °C for 1 h. The reaction mixture was then filtered through a pad of celite and concentrated to obtain a thick oil. The residue was taken up in EtOAc (200 mL) and sat. NaHCO₃ (100 mL). The organic layer was separated, washed with H₂O (2x 50 mL), dried over Na₂SO₄, and concentrated. Column chromatography (silica, 10-30% EtOAc/hexanes) provided the desired amine (844 mg, 92%). The amine (42 mg, 0.13 mmol) in CH₂Cl₂ (0.5 mL) was treated with 9.7 μL (0.13 mmol, 1.0 eq) of methanesulfonyl chloride and 34.9 μL (0.25 mmol, 2 eq) of triethylamine and the reaction mixture stirred at 25 °C for 1 h. EtOAc (20 mL) and sat. NaHCO₃ (10 mL) were added. The organic layer was separated and washed with H₂O (10 mL), brine (20 mL), dried over Na₂SO₄, and concentrated. The crude oil was dissolved in 2 mL of CH₂Cl₂ and treated with 0.5 mL of trifluoroacetic acid. The reaction mixture was stirred at 25 °C for 1 h before all volatiles were removed. The crude oil was dissolved in absolute EtOH (1.0 mL) and treated

with 40 mg (0.1 mmol) of 5-methanesulfonyl-1-oxiranylmethyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine and 200 μ L of triethylamine. The reaction mixture was refluxed at 85 °C for 4 h and then the solvent was removed. Preparative TLC (silica, 7% MeOH/CH₂Cl₂) provided

5 35 mg (49% over 3 steps) of the title compound. TLC (silica, 5% MeOH/CH₂Cl₂): R_f = 0.30. MS (electrospray): exact mass calculated for C₃₀H₃₇F₃N₆O₇S₂, 714.21; m/z found, 715.2 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃): 8.11 (s, 1H), 7.74-7.59 (m, 5H), 7.30 (d, J = 8.1 Hz, 1H), 7.21 (t, J = 8.2 Hz, 1H), 4.62-4.49 (m, 2H), 4.25-3.99 (m, 3H), 3.90 (s, 3H), 3.80-3.57 (m, 3H),

10 3.53-3.27 (m, 2H), 3.14-2.78 (m, 4H), 3.05 (s, 3H), 2.86 (s, 3H), 2.76-2.65 (m, 2H), 2.61-2.20 (m, 4H).

EXAMPLE 22



15 1-{3-[4-(1,1-Dioxo-1H-1H-6-benzo[d]isothiazol-3-yl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine.

20

A. 3-Piperazin-1-yl-benzo[d]isothiazole 1,1-dioxide.

POCl₃ (10.2 mL, 109.2 mmol) was added to saccharin (5.0 g, 27.3 mmol) and heated at 120 °C for 20 h. The excess reagent was removed in a rotary evaporator and water (50 mL) was added to the residue to form a precipitate.

25 The solid was filtered, washed with water (2 x 20 mL), and dried. A portion of the above crude material (2.0 g, 9.95 mmol) and piperazine (4.28 g, 49.75 mmol) was taken in dioxane (10 mL), and heated at 100 °C for 24 h. The reaction was allowed to cool to room temperature and poured into ice water (50 g), and neutralized by addition of 10% aqueous NaOH. The mixture was

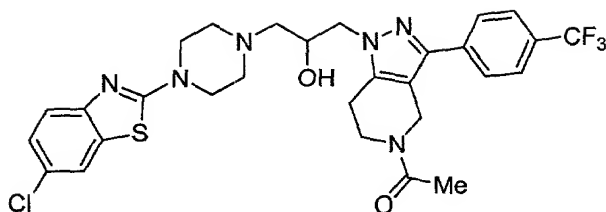
extracted with CH_2Cl_2 (3 x 25 mL) and the combined organic extracts were washed with brine, dried (Na_2SO_4) and concentrated. MPLC (silica, 5-20% $\text{MeOH}/\text{CH}_2\text{Cl}_2$) afforded the piperazinyl derivative (0.07 g, 4.2%). MS

(electrospray): exact mass calculated for $\text{C}_{11}\text{H}_{13}\text{N}_3\text{O}_2\text{S}$, 251.07; m/z found, 252.1 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3): 7.72 (dd, $J = 0.8, 7.4$ Hz, 1H), 7.64 (d, $J = 7.8$ Hz, 1H), 7.49 (dt, $J = 0.8, 7.4$ Hz, 1H), 7.43 (dt, $J = 1.2, 7.8$ Hz, 1H), 3.80 (s, 4H), 2.85 (br t, $J = 5.0$ Hz, 4H), 2.07 (br s, 1H). ^{13}C NMR (100 MHz, CDCl_3): 160.8, 145.3, 133.3, 133.0, 128.5, 125.9, 123.2, 49.8, 46.3.

10 B. 1-{3-[4-(1,1-Dioxo-1H-1*b*-benzo[d]isothiazol-3-yl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine.

To a stirred solution of 3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propionaldehyde (0.040 g, 0.13 mmol) and 3-piperazin-1-yl-benzo[d]isothiazole 1,1-dioxide (0.050 g, 0.21 mmol) in CH_2Cl_2 (0.5 mL), glacial AcOH (12 μL , 0.21 mmol) was added and stirred for 15 min at room temperature. $\text{NaBH}(\text{OAc})_3$ (0.058 g, 0.27 mmol) was added and stirred under nitrogen overnight. Saturated NaHCO_3 (0.5 mL) was then added and stirred for 15 min. The layers separated and the aqueous layer was extracted with CH_2Cl_2 (0.5 mL). MPLC (silica, 2-15% $\text{MeOH}/\text{CH}_2\text{Cl}_2$) afforded the desired product as a white solid (0.048 g, 76%). TLC (silica, 12% $\text{MeOH}/\text{CH}_2\text{Cl}_2$): $R_f = 0.50$. MS (electrospray): exact mass calculated for $\text{C}_{28}\text{H}_{31}\text{F}_3\text{N}_6\text{O}_4\text{S}_2$, 636.18; m/z found, 637.2 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3): 7.94 (dd, $J = 0.8, 7.6$ Hz, 1H), 7.86 (d, $J = 7.8$ Hz, 1H), 7.75 (d, $J = 8.3$ Hz, 2H), 7.73-7.63 (m, 2H), 7.68 (d, $J = 8.3$ Hz, 2H), 4.57 (s, 2H), 4.17 (t, $J = 6.9$ Hz, 2H), 4.04 (br s, 4H), 3.69 (t, $J = 5.7$ Hz, 2H), 2.94 (s, 3H), 2.92 (t, $J = 6.2$ Hz, 2H), 2.62 (t, $J = 5.0$ Hz, 4H), 2.44 (t, $J = 6.6$ Hz, 2H), 2.13 (q, $J = 6.6$ Hz, 2H).

EXAMPLE 23



- 1-[1-{3-[4-(6-Chloro-benzothiazol-2-yl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

A. 6-Chloro-2-piperazin-1-yl-benzothiazole.

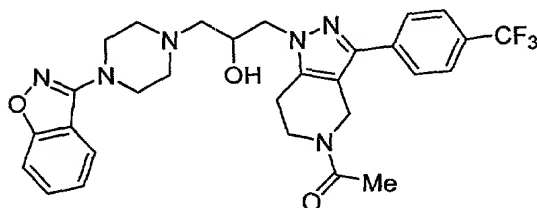
- To a stirred solution of 1.07 g (5.24 mmol) of 2,6-dichlorobenzothiazole in dry DMF (25 mL) was added 2.4 g of potassium carbonate (15.7 mmol) and 0.5 g of piperazine (5.8 mmol). The mixture was stirred at room temperature for 4 h. When the reaction was complete it was partitioned between EtOAc (150 mL) and water (50 mL) and separated. The aqueous layer was extracted with EtOAc (2 x 100 mL). The combined organic layers were then washed with water (2 x 25 mL), brine, dried over Na₂SO₄, and the solvent was removed under reduced pressure to give 1.33 g (100%) of desired product as a white solid. MS (electrospray): exact mass calculated for C₁₁H₁₂ClN₃S, 253.04; *m/z* found, 254.0 [M+H]⁺.

- B. 1-[1-{3-[4-(6-Chloro-benzothiazol-2-yl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

- To a stirred solution of 144 mg (0.39 mmol) of 1-[1-oxiranylmethyl-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone in 4 mL of EtOH was added 100 mg (0.39 mmol) 6-chloro-2-piperazin-1-yl-benzothiazole. The solution was heated to 60 °C overnight. The solvent was then removed by rotary evaporation and the crude product was purified by column chromatography (silica, 0-10% MeOH/EtOAc) to afford 220 mg (90%) of a white solid. MS (electrospray): exact mass calculated for C₂₉H₃₀ClF₃N₆O₂S:

618.18; m/z found, 619.2 $[M+H]^+$. HPLC (reverse phase conditions 40-90%): t_R = 8.27 min. 1H NMR ($CDCl_3$, 400 MHz, a mixture of amide rotamers): 7.70 (d, J = 8.34 Hz, 1H), 7.62 (m, 2H), 7.57 (d, J = 8.59 Hz, 1H), 7.48 (d, J = 2.53 Hz, 1H), 7.36 (d, J = 8.59 Hz, 1H), 7.16 (dd, J = 8.59, 2.53 Hz, 1H), 4.80 and 4.68 (A and B of AB quartet, J = 15.92 Hz, 1H), 4.58 (s, 1H), 4.18-4.08 (m, 2H), 4.01-3.89 (m, 2H), 3.85-3.60 (m, 2H), 3.59-3.47 (m, 4H), 2.94-2.75 (m, 2H), 2.72-2.62 (m, 2H), 2.55-2.47 (m, 2H), 2.46-2.39 (m, 2H), 2.13 (s, 1.5H), 2.08 (s, 1.5H).

EXAMPLE 24



1-[1-[3-(4-Benzo[d]isoxazol-3-yl)-piperazin-1-yl]-2-hydroxy-propyl]-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

A. 4-Benzo[d]isoxazol-3-yl-piperazine-1-carboxylic acid tert-butyl ester.

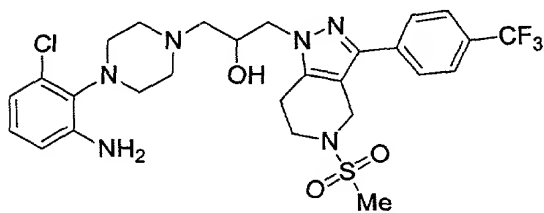
To a stirred solution of 100 mg (0.65 mmol) of 3-chloro-1,2-benzisoxazole in pyridine (1 mL) was added 145 mg of piperazine-1-carboxylic acid tert-butyl ester (0.78 mmol) and 0.18 mL of DBU (0.78 mmol). The mixture was stirred at 100 °C overnight and then partitioned between EtOAc (50 mL) and water (20 mL) and separated. The aqueous layer was extracted with EtOAc (2 x 30 mL). The combined organic layers were then washed with water (25 mL), brine, dried over Na_2SO_4 , and the solvent was removed under reduced pressure to give crude product. Purification by column chromatography (silica, 60-100% CH_2Cl_2 /hexanes) gave 82 mg (42%) of the desired product as a light yellow solid. MS (electrospray): exact mass calculated for $C_{16}H_{21}N_3O_3$, 303.16; m/z found, 326.1 $[M+Na]^+$. 1H NMR ($CDCl_3$, 400 MHz): 7.68 (dt, J = 8.02, 0.98

Hz, 1H), 7.52-7.44 (m, 2H), 7.24 (ddd, $J = 8.42, 6.46, 1.57$ Hz, 1H), 3.66-3.61 (m, 4H), 3.56-3.49 (m, 4H), 1.49 (s, 9H).

B. 1-[1-[3-(4-Benzo[d]isoxazol-3-yl-piperazin-1-yl)-2-hydroxy-propyl]-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone.

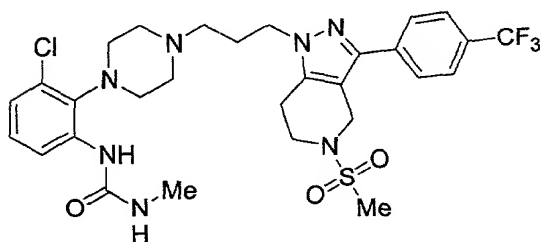
A solution of 82 mg (0.27 mmol) of 4-benzo[d]isoxazol-3-yl-piperazine-1-carboxylic acid tert-butyl ester in 2 mL of CH_2Cl_2 was treated with trifluoroacetic acid (0.5 mL) at room temperature overnight. The solvent was then removed and the crude product dissolved in EtOH and stirred over 100 mg of sodium bicarbonate for 1 h, the solid was then filtered off and the filtrate concentrated. The crude piperazine was then dissolved in 4 mL EtOH and treated with 100 mg (0.27 mmol) of 1-[1-oxiranylmethyl-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone. The solution was heated to 60 °C overnight. The solvent was then removed by rotary evaporation and the crude product was purified by column chromatography (silica, 0-10% MeOH/EtOAc) to afford 105 mg (68%) of a white solid. MS (electrospray), exact mass calculated for $\text{C}_{29}\text{H}_{31}\text{F}_3\text{N}_6\text{O}_3$, 568.24; m/z found, 569.2 $[\text{M}+\text{H}]^+$. ^1H NMR (CDCl_3 , 400 MHz, a mixture of amide rotamers): 7.77 (d, $J = 8.41$ Hz, 1H), 7.69 (m, 2H), 7.67-7.62 (m, 2H), 7.50-7.44 (m, 1H), 7.45-7.42 (m, 1H), 7.23-7.18 (m, 1H), 4.93 (br m, 1H), 4.87 and 4.75 (A and B of AB quartet, $J = 15.65$ Hz, 1H), 4.65 (br s, 1H), 4.27-4.15 (m, 2.3H), 4.09-3.95 (m, 1.7H), 3.91-3.82 (m, 0.7H), 3.81-3.66 (m, 1.3H), 3.62-3.49 (m, 4H), 3.01-2.85 (m, 1.5H), 2.85-2.74 (m, 2.5H), 2.71-2.60 (m, 2H), 2.58-2.45 (m, 2H), 2.20 (s, 1.5H), 2.15 (s, 1.5H).

EXAMPLE 25



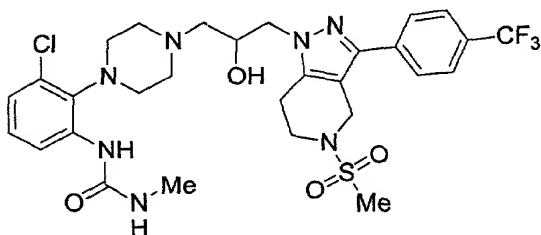
1-[4-(2-Amino-6-chloro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol.

5 EXAMPLE 26



10 1-[3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-3-methyl-urea.

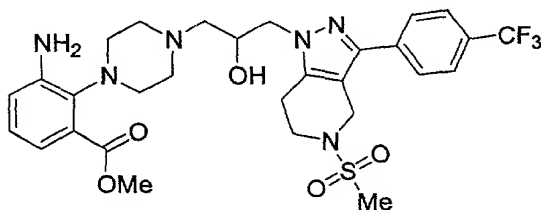
EXAMPLE 27



15 1-[3-Chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-3-methyl-urea.

EXAMPLE 28

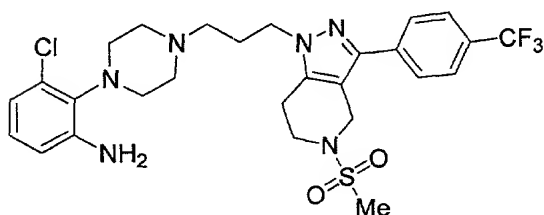
20



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3-Amino-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzoic acid methyl ester.

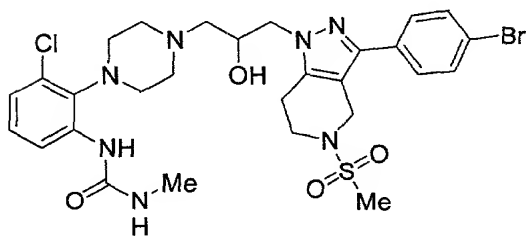
5 EXAMPLE 29



3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenylamine.

10

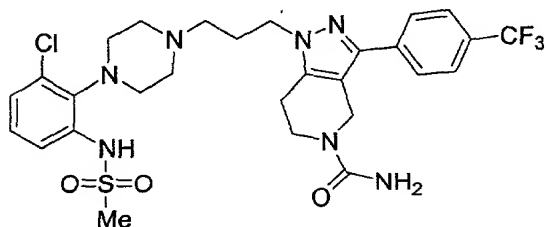
EXAMPLE 30



15 1-[2-(4-{3-[3-(4-Bromo-phenyl)-5-methanesulfonyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-3-chloro-phenyl]-3-methyl-urea.

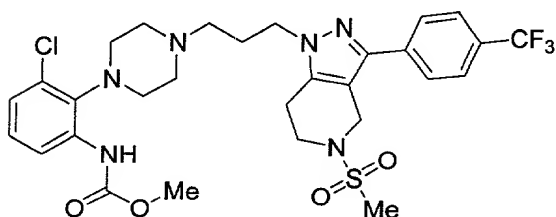
EXAMPLE 31

20



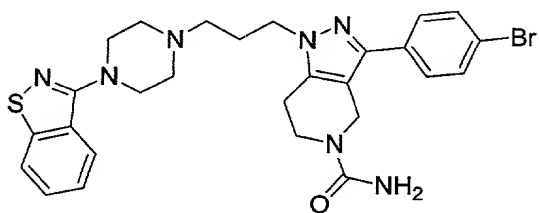
1-{3-[4-(2-Chloro-6-methanesulfonylamino-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide.

5 EXAMPLE 32



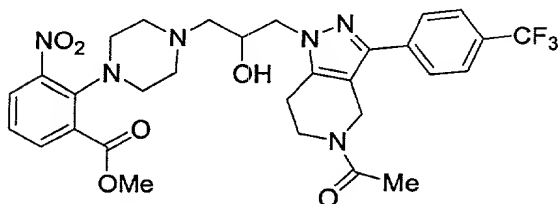
[3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-carbamic acid methyl ester.

EXAMPLE 33



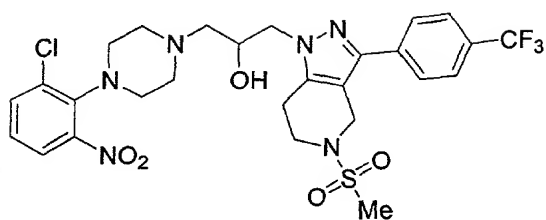
1-[3-(4-Benzo[d]isothiazol-3-yl-piperazin-1-yl)-propyl]-3-(4-bromo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide.

EXAMPLE 34



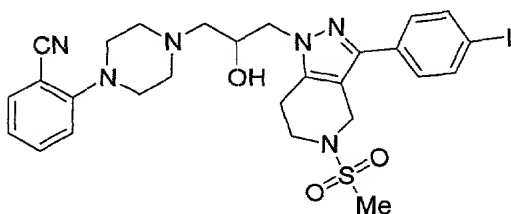
2-(4-{3-[5-Acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-3-nitro-benzoic acid methyl ester.

EXAMPLE 35



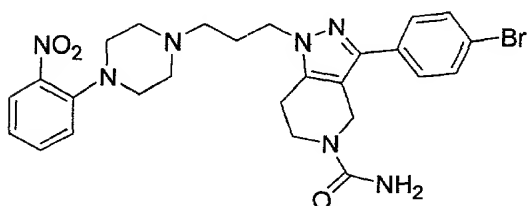
- 5 1-[4-(2-Chloro-6-nitro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol.

EXAMPLE 36



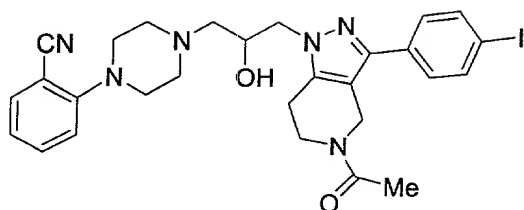
- 10 2-(4-{2-Hydroxy-3-[3-(4-iodo-phenyl)-5-methanesulfonyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzonitrile.

EXAMPLE 37



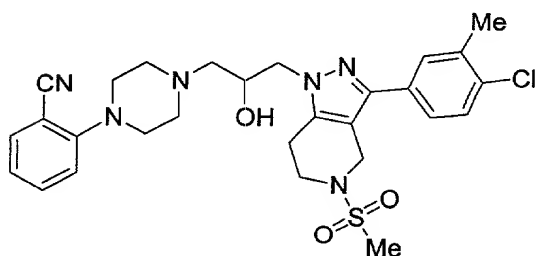
- 15 3-(4-Bromo-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide.

EXAMPLE 38



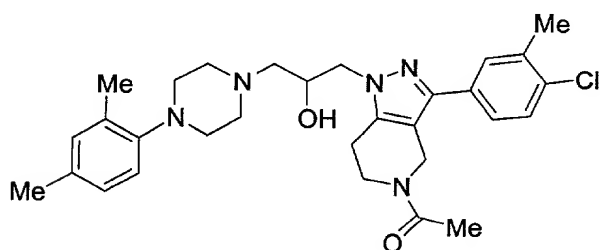
2-(4-{3-[5-Acetyl-3-(4-iodo-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile.

EXAMPLE 39



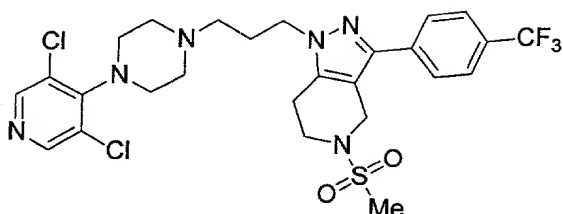
2-(4-{3-[3-(4-Chloro-3-methyl-phenyl)-5-methanesulfonyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile.

EXAMPLE 40



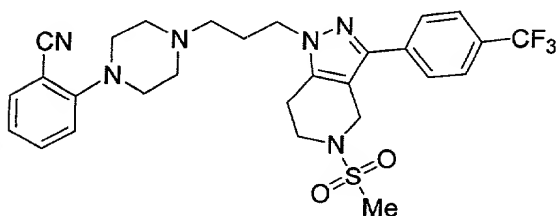
1-(3-(4-Chloro-3-methyl-phenyl)-1-{3-[4-(2,4-dimethyl-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone.

EXAMPLE 41



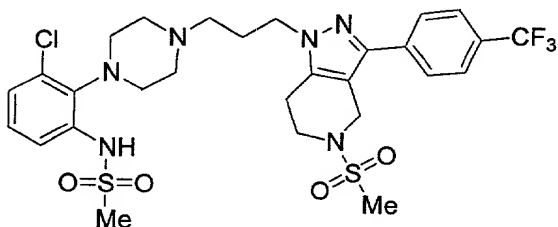
1-{3-[4-(3,5-Dichloro-pyridin-4-yl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-3-
5 (4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine.

EXAMPLE 42



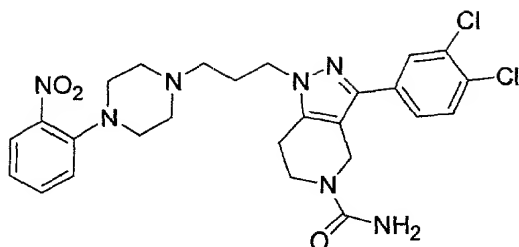
10 2-(4-{3-[5-Methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-
pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzonitrile.

EXAMPLE 43



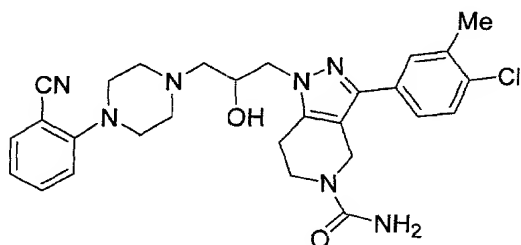
15 N-[3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-
tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-
methanesulfonamide.

EXAMPLE 44



3-(3,4-Dichloro-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-
5 tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide.

EXAMPLE 45



3-(4-Chloro-3-methyl-phenyl)-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-
10 hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid
amide.

EXAMPLE 46

Cathepsin S Inhibition Assay.

15

Recombinant human cathepsin S (CatS) was expressed in the
baculovirus system and purified in one step with a thiopropyl-sepharose
column. 10-L yielded ~700 mg of CatS and N-terminal sequencing confirmed
identity. The assay is run in 100 mM sodium acetate pH 5.0 containing 1 mM
20 DTT and 100 mM NaCl. The substrate for the assay is

(Aedens)EKARVLAEAA(Dabcyl)K-amide

The K_m for the substrate is around 5 μ M but the presence of substrate inhibition
25 makes kinetic analysis difficult. With 20 μ M substrate the assay rate is linear

over the range of 1-8 ng CatS in 100 μ l reaction. Using 2 ng/well of CatS, the production of product is linear and yields ~7-fold signal after 20 min with only 20% loss of substrate. Primary assays are run by quenching the reaction after 20 min with 0.1% SDS and then measuring the fluorescence. For other

5 assays, measurements are taken every min for 20 min. The rate is calculated from the slope of the increase and the percent inhibition is calculated from this (See Tables 1, 2 and 3 below).

Table 1

EXAMPLE	IC ₅₀ (μM)
1	0.89
2	1.22
3	0.84
4	0.51
5	0.36
6	0.30
7	6.60
8	0.89
9	1.14
10	0.05
11	0.03
12	0.98
13	0.77
14	0.25
15	0.12
16	0.06
17	0.08
18	0.14
19	0.06
20	0.17
21	0.07
22	2.15
23	1.10
24	0.47

Table 2

EXAMPLE	IC ₅₀ (μM)
25	0.04
26	0.04
27	0.04
28	0.07
29	0.07
30	0.08
31	0.10
32	0.10
33	0.10
34	0.11
35	0.12
36	0.12
37	0.12
38	0.12
39	0.13
40	0.13
41	0.13
42	0.13
43	0.13
44	0.13
45	0.13

EXAMPLE 101

1-(3-(4-Chloro-phenyl)-1-{4-[4-(2-methoxy-phenyl)-piperazin-1-yl]-butyl}-
1,4,6,7-tetrahydro-pyrazolo [4,3-c]pyridin-5-yl)-ethanone

5 EXAMPLE 102

1-[1-(3-{4-[Bis-(4-fluoro-phenyl)-methyl]-piperazin-1-yl}-2-hydroxy-propyl)-3-(4-
chloro-phenyl)-1,4, 6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 103

10 1-(3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-
propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 104

15 1-(3-(4-Chloro-phenyl)-1-{3-[4-(2-chloro-phenyl)-piperazin-1-yl]-2-hydroxy-
propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 105

20 1-(3-(4-Chloro-phenyl)-1-{3-[4-(3-chloro-phenyl)-piperazin-1-yl]-2-hydroxy-
propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 106

1-(3-(4-Chloro-phenyl)-1-{3-[4-(4-chloro-phenyl)-piperazin-1-yl]-2-hydroxy-
propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

25 EXAMPLE 107

1-(3-(4-Chloro-phenyl)-1-{3-[4-(2-fluoro-phenyl)-piperazin-1-yl]-2-hydroxy-
propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 108

30 1-(3-(4-Chloro-phenyl)-1-{3-[4-(4-fluoro-phenyl)-piperazin-1-yl]-2-hydroxy-
propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 109

1-(3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(3-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

5 EXAMPLE 110

1-(3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(4-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 111

10 1-{3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-phenyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

EXAMPLE 112

15 1-[1-[3-(4-Benzhydryl-piperazin-1-yl)-2-hydroxy-propyl]-3-(4-chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 113

1-[3-(4-Chloro-phenyl)-1-(3-{4-[(4-chloro-phenyl)-phenyl-methyl]-piperazin-1-yl}-2-hydroxy-propyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 114

1-(3-(4-Chloro-phenyl)-1-{3-[4-(9H-fluoren-9-yl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

25 EXAMPLE 115

1-[1-[3-(4-Benzyl-piperazin-1-yl)-2-hydroxy-propyl]-3-(4-chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 116

30 3-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-1-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propan-1-one

EXAMPLE 117

1-[1-{2-Hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

5 EXAMPLE 118

1-(3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(2-trifluoromethyl-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 119

10 1-(3-(4-Fluoro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 120

15 4-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-1-[4-(2-methoxy-phenyl)-piperazin-1-yl]-butan-1-one

EXAMPLE 121

20 1-(1-{2-Hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-3-p-tolyl-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 122

1-(3-(4-Chloro-phenyl)-1-{3-[4-(3,4-dichloro-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

25 EXAMPLE 123

1-{3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-pyridin-2-yl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

EXAMPLE 124

30 1-(3-Biphenyl-4-yl-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 125

1-(1-{2-Hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-3-phenyl-1,4,6,7-tetrahydro-pyrazolo [4,3-c]pyridin-5-yl)-ethanone

5 EXAMPLE 126

1-[1-{2-Hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-3-(4-methoxy-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 127

10 1-[1-[2-Hydroxy-3-(4-pyridin-4-yl-piperazin-1-yl)-propyl]-3-(4-methoxy-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 129

15 1-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-one

EXAMPLE 130

20 3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester

EXAMPLE 131

25 1-(1-{2-Hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-3-naphthalen-2-yl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 132

1-(3-(4-tert-Butyl-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

30 EXAMPLE 133

1-(3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-butan-1-one

EXAMPLE 134

1-(3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-2,2-dimethyl-propan-1-one

5

EXAMPLE 136

(3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-(4-methoxy-phenyl)-methanone

10

EXAMPLE 137

3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide

15

EXAMPLE 138

1-[3-(4-Chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propan-2-ol

EXAMPLE 139

1-(3-(3,4-Dichloro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

20

EXAMPLE 140

1-[1-{2-Hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

25

EXAMPLE 141

1-[1-{2-Hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-3-(4-nitro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

30

EXAMPLE 142

1-(3-(4-Chloro-phenyl)-1-{3-[4-(2,4-difluoro-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 143

2-(4-{3-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

5

EXAMPLE 144

4-{3-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-3,4,5,6-tetrahydro-2H-[1,2']bipyrazinyl-3'-carbonitrile

10 EXAMPLE 145

1-(3-(4-Chloro-phenyl)-1-{3-[4-(2,3-dimethyl-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 146

15 1-(3-(4-Chloro-phenyl)-1-{3-[4-(2,4-dimethyl-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 147

20 1-(3-(4-Chloro-phenyl)-1-{3-[4-(2,5-dimethyl-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 148

25 1-{3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(3-methyl-4-p-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

EXAMPLE 149

1-{3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(3-methyl-4-m-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

30 EXAMPLE 150

1-(3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(4-trifluoromethyl-pyridin-2-yl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 151

1-(3-(4-Chloro-phenyl)-1-{3-[4-(3-chloro-5-trifluoromethyl-pyridin-2-yl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

5

EXAMPLE 152

1-(3-(4-Chloro-phenyl)-1-{3-[4-(3,5-dichloro-pyridin-4-yl)-piperazin-1-yl]-2-hydroxy-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

10 EXAMPLE 153

1-(3-(4-Chloro-phenyl)-1-{4-[4-(2-methoxy-phenyl)-piperazin-1-yl]-but-2-enyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 154

15 4-(5-Acetyl-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridin-3-yl)-benzonitrile

EXAMPLE 155

20 1-{3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(2,3,5,6-tetrahydro-[1,2']bipyrazinyl-4-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 156

1-{3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-pyrimidin-2-yl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

25

EXAMPLE 157

1-(3-(2,4-Bis-trifluoromethyl-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

30 EXAMPLE 158

1-(3-(2,4-Dichloro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 159

2-(4-{3-[3-(4-Chloro-phenyl)-5,6-dihydro-4H-cyclopentapyrazol-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

5 EXAMPLE 160

2-(4-{3-[3-(4-Chloro-phenyl)-5,6-dihydro-4H-cyclopentapyrazol-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-phenol

EXAMPLE 161

10 1-(3-(4-Bromo-phenyl)-1-[2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 162

15 1-{3-(4-Chloro-phenyl)-1-[2-(2-methyl-allyloxy)-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

EXAMPLE 163

20 1-[1-[2-Benzoyloxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 164

Acetic acid 1-[5-acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-ylmethyl]- 2-(4-o-tolyl-piperazin-1-yl)-ethyl ester

25 EXAMPLE 165

Morpholine-4-carboxylic acid 1-[5-acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-ylmethyl]-2-(4-o-tolyl-piperazin-1-yl)-ethyl ester

EXAMPLE 166

30 Benzoic acid 1-[5-acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-ylmethyl] -2-(4-o-tolyl-piperazin-1-yl)-ethyl ester

EXAMPLE 167

Benzoyl-carbamic acid 1-[5-acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-ylmethyl]-2-(4-o-tolyl-piperazin-1-yl)-ethyl ester

5 EXAMPLE 168

1-[3-(4-Chloro-phenyl)-pyrazol-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol

EXAMPLE 169

10 1-(3-(3-Chloro-phenyl)-1-{2-hydroxy-3-[4-(2-hydroxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 170

15 2-(4-{3-[5-Acetyl-3-(3-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 171

tert-Butyl-carbamic acid 1-[5-acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-ylmethyl]-2-(4-o-tolyl-piperazin-1-yl)-ethyl ester

20 EXAMPLE 172

Carbonic acid 1-[5-acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-ylmethyl]-2-(4-o-tolyl-piperazin-1-yl)-ethyl ester methyl ester

EXAMPLE 173

25 1-(3-(4-Chloro-phenyl)-1-{4-[4-(2-hydroxy-phenyl)-piperazin-1-yl]-but-2-enyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 174

30 2-(4-{4-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-but-2-enyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 175

1-(3-(4-Chloro-phenyl)-1-{4-[4-(2-methoxy-phenyl)-piperazin-1-yl]-but-2-enyl}-
1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

5 EXAMPLE 176

1-(3-(4-Chloro-phenyl)-1-{3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-
1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 177

10 1-(3-(4-Chloro-phenyl)-1-{5-[4-(2-methoxy-phenyl)-piperazin-1-yl]-pentyl}-
1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 178

15 1-(3-(4-Chloro-phenyl)-1-{6-[4-(2-methoxy-phenyl)-piperazin-1-yl]-hexyl}-
1,4,6,7-tetrahydro-pyrazolo [4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 179

20 2-[1-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-
ylmethyl]-2-(4-o-tolyl-piperazin-1-yl)-ethoxy]-acetamide

EXAMPLE 180

[1-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-
ylmethyl]-2-(4-o-tolyl-piperazin-1-yl)-ethoxy]-acetic acid

25 EXAMPLE 181

[1-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-
ylmethyl]-2-(4-o-tolyl-piperazin-1-yl)-ethoxy]-acetonitrile

EXAMPLE 182

30 1-[1-{3-[4-(2-Bromo-benzenesulfonyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-
chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 183

3-(5-(4-Chloro-phenyl)-2-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-2H-pyrazol-3-yl) -propionic acid methyl ester

5 EXAMPLE 184

2-(4-{3-[3-(4-Chloro-phenyl)-4,5,6,7-tetrahydro-indazol-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 185

10 2-(4-{3-[3-(4-Chloro-phenyl)-4,5,6,7-tetrahydro-indazol-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-phenol

EXAMPLE 186

15 3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid dimethylamide

EXAMPLE 187

20 1-[1-[2-Azido-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo [4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 188

1-[1-[2-Amino-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo [4,3-c]pyridin-5-yl]-ethanone

25 EXAMPLE 189

1-{3-(4-Chloro-phenyl)-1-[2-methylamino-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

EXAMPLE 190

30 3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide

EXAMPLE 191

3-(4-Chloro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-indazol-5-one ethylene ketal

5 EXAMPLE 192

1-[2-Hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester

EXAMPLE 193

10 1-(3-(4-Chloro-3-methyl-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 194

15 2-(4-{3-[5-Acetyl-3-(4-chloro-3-methyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 195

20 1-[1-{3-[4-(2-Chloro-benzenesulfonyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 196

1-(3-(4-Chloro-2-fluoro-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

25 EXAMPLE 197

2-(4-{3-[5-Acetyl-3-(4-chloro-2-fluoro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 198

30 1-[3-(4-Chloro-phenyl)-5-methyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol

EXAMPLE 199

1-[3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-2-phenyl-ethanone

5 EXAMPLE 200

1-[3-(4-Chloro-phenyl)-5-methanesulfonyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol

EXAMPLE 201

10 1-[1-[3-[4-(2-Amino-phenyl)-piperazin-1-yl]-2-hydroxy-propyl]-3-(4-chloro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 202

15 N-[2-(4-{3-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-phenyl]-methanesulfonamide

EXAMPLE 203

20 N-[2-(4-{3-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-phenyl]-acetamide

EXAMPLE 204

1-[2-(4-{3-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-phenyl]-3-isopropyl-urea

25 EXAMPLE 205

3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid methylamide

EXAMPLE 206

30 3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid hydrazide

EXAMPLE 207

2-(4-{3-[5-Acetyl-3-(4-phenoxy-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

5 EXAMPLE 208

3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid phenethyl-amide

EXAMPLE 209

10 3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid (4-methoxy-phenyl)-amide

EXAMPLE 210

15 3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carbothioic acid methylamide

EXAMPLE 211

20 2-(4-{3-[5-Acetyl-3-(4-chloro-3-nitro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 212

1-[2-Hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4, 3-c]pyridine-5-carboxylic acid ethylamide

25 EXAMPLE 213

N-(5-{5-Acetyl-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridin-3-yl}-2-chloro-phenyl)-methanesulfonamide

30 EXAMPLE 214

1-{3-(4-Chloro-phenyl)-1-[2-[(1-ethyl-pyrrolidin-2-ylmethyl)-amino]-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

EXAMPLE 215

2-(4-{3-[5-Acetyl-3-(4-trifluoromethylsulfanyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

5 EXAMPLE 216

2-(4-{3-[5-Acetyl-3-(3-amino-4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 217

10 3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid isopropylamide

EXAMPLE 218

15 3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid phenylamide

EXAMPLE 219

20 1-[3-(4-Chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol

EXAMPLE 220

1-[3-(4-Iodo-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol

25 EXAMPLE 221

2-(4-{3-[5-Acetyl-3-(4-methanesulfonyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 222

30 1-[1-{2-Hydroxy-3-[4-(2-hydroxy-phenyl)-piperazin-1-yl]-propyl}-3-(4-methanesulfonyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 223

11-[3-(4-Iodo-phenyl)-5-methanesulfonyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol

5 EXAMPLE 224

1-[2-Hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4, 3-c]pyridine-5-carboxylic acid amide

EXAMPLE 225

10 1-[2-Hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid methyl ester

EXAMPLE 226

15 1-[2-Hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid methylanide

EXAMPLE 227

20 N-[5-(5-Acetyl-1-[3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl]-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridin-3-yl)-2-chloro-phenyl]-methanesulfonamide

EXAMPLE 228

25 1-(5-{5-Acetyl-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridin-3-yl}-2-chloro-phenyl)-3-ethyl-urea

EXAMPLE 229

1-[5-(5-Acetyl-1-[3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl]-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridin-3-yl)-2-chloro-phenyl]-3-ethyl-urea

30 EXAMPLE 230

N-(5-{5-Acetyl-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridin-3-yl}-2-chloro-phenyl)-acetamide

EXAMPLE 231

Acetic acid 2-[5-acetyl-3-(3-amino-4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-1-[4-(2-cyano-phenyl)-piperazin-1-ylmethyl]-ethyl ester

5

EXAMPLE 232

N-[5-(5-Acetyl-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridin-3-yl)-2-chloro-phenyl]-acetamide

10 EXAMPLE 233

N-[2-[5-Acetyl-3-(4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-1-(4-o-tolyl-piperazin-1-ylmethyl)-ethyl]-methanesulfonamide

EXAMPLE 234

15 1-{3-(4-Chloro-phenyl)-1-[2-(2-pyridin-2-yl-ethylamino)-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

EXAMPLE 235

20 1-{3-(4-Chloro-phenyl)-1-[2-(2-dimethylamino-ethylamino)-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4, 6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl}-ethanone

EXAMPLE 236

25 Carbonic acid 2-[5-acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-1-(4-o-tolyl-piperazin-1-ylmethyl)-ethyl ester methyl ester

EXAMPLE 237

Carbamic acid 2-[5-acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-1-(4-o-tolyl-piperazin-1-ylmethyl)-ethyl ester

30

EXAMPLE 238

3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-indazol-5-one

EXAMPLE 239

3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-4,5,6,7-tetrahydro-1H-indazol-5-ol

5

EXAMPLE 240

3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-indazol-5-one oxime

10 EXAMPLE 241

1-[5-Ethanesulfonyl-3-(4-iodo-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol

EXAMPLE 242

15 1-[2-Hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4, 3-c]pyridine-5-carboxylic acid methyl ester

EXAMPLE 243

20 1-[5-(4-Chloro-benzenesulfonyl)-3-(4-iodo-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]- 3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol

EXAMPLE 244

25 1-{3-[4-(2-Cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid methylamide

EXAMPLE 245

1-[3-(4-Iodo-phenyl)-5-(propane-2-sulfonyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-3-(4-o-tolyl-piperazin-1-yl)-propan-2-ol

30 EXAMPLE 246

1-{3-[4-(2-Cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-iodo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carbonitrile

EXAMPLE 247

4-{3-[5-Acetyl-3-(4-chloro-3-methyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazine-1-carboxylic acid o-tolylamide

5 EXAMPLE 248

4-{3-[5-Acetyl-3-(4-chloro-3-methyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazine-1-carboxylic acid (2-methoxy-phenyl)-amide

10 EXAMPLE 249

2-(4-{3-[5-Acetyl-3-(3-chloro-4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 250

15 2-(4-{3-[5-Acetyl-3-(3-fluoro-4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 251

20 2-(4-{3-[5-Acetyl-3-(4-chloro-3-methyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-ylmethyl)-benzonitrile

EXAMPLE 252

25 1-(3-(4-Chloro-3-methyl-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-benzyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 253

2-(4-{3-[5-Acetyl-3-(4-bromo-3-methyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

30 EXAMPLE 254

3-(4-Chloro-phenyl)-1-[2-hydroxy-3-(4-o-tolyl-piperazin-1-yl)-propyl]-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxamidine

EXAMPLE 255

2-(4-{3-[5-Acetyl-3-(3,4-dichloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

5 EXAMPLE 256

2-(4-{3-[5-Acetyl-3-(3,4-difluoro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 257

10 2-(4-{3-[5-Acetyl-3-(3,5-dichloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 258

15 2-{4-[3-[5-Acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-(2-morpholin-4-yl-ethoxy)-propyl]-piperazin-1-yl}-benzonitrile

EXAMPLE 259

20 2-(4-{2-Hydroxy-3-[3-(4-iodo-phenyl)-5-trifluoromethanesulfonyl-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 260

2-(4-{3-[5-Acetyl-3-(3-chloro-4-methyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

25 EXAMPLE 261

N-[4-(5-Acetyl-1-{3-[4-(2-cyano-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridin-3-yl)-phenyl]-acetamide

EXAMPLE 262

30 2-(4-{3-[5-Acetyl-3-(4-bromo-3-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 263

1-(3-(3-Chloro-4-methyl-phenyl)-1-{2-hydroxy-3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

5 EXAMPLE 264

1-[1-{3-[4-(2-Azido-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-bromo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 265

10 2-(4-{3-[5-Acetyl-3-(3-azido-4-chloro-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 266

15 5-Methanesulfonyl-1-[3-(4-o-tolyl-piperazin-1-yl)-propyl]-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

EXAMPLE 267

20 5-Methanesulfonyl-1-{3-[4-(2-methoxy-phenyl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

EXAMPLE 268

1-[1-{2-Hydroxy-3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-3-(4-nitro-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

25 EXAMPLE 269

3-(4-Bromo-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester

EXAMPLE 270

30 3-(4-Bromo-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

EXAMPLE 271

1-(3-(4-Bromo-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

5 EXAMPLE 272

3-(4-Bromo-phenyl)-5-methanesulfonyl-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

EXAMPLE 273

10 3-(3,4-Dichloro-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester

EXAMPLE 274

15 3-(4-Bromo-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-sulfonic acid amide

EXAMPLE 275

20 1-(3-(3,4-Dichloro-phenyl)-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl)-ethanone

EXAMPLE 276

3-(3,4-Dichloro-phenyl)-5-methanesulfonyl-1-{3-[4-(2-nitro-phenyl)-piperazin-1-yl]-propyl}-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

25 EXAMPLE 277

3-(4-Bromo-phenyl)-1-{3-[4-(1,1-dioxo-1H-116-benzo[d]isothiazol-3-yl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

30 EXAMPLE 278

1-[1-[3-(4-Benzo[d]isothiazol-3-yl-piperazin-1-yl)-propyl]-3-(4-bromo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 279

1-[3-(4-Benzo[d]isothiazol-3-yl-piperazin-1-yl)-propyl]-3-(4-bromo-phenyl)-5-methanesulfonyl-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

5 EXAMPLE 280

1-[3-(4-Benzo[d]isothiazol-3-yl-piperazin-1-yl)-propyl]-3-(4-bromo-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid tert-butyl ester

EXAMPLE 281

10 1-[3-(4-Benzo[d]isothiazol-3-yl-piperazin-1-yl)-propyl]-3-(4-bromo-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

EXAMPLE 282

15 1-[3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-3-methyl-urea

EXAMPLE 283

20 [3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-urea

EXAMPLE 284

25 [3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-carbamic acid methyl ester

EXAMPLE 285

30 1-[3-Chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-3-methyl-urea

EXAMPLE 286

N-[3-Chloro-2-(4-{3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-methanesulfonamide

5

EXAMPLE 287

1-[4-(2,6-Dimethyl-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol

10

EXAMPLE 288

1-[1-{3-[4-(2,6-Dimethyl-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

15

EXAMPLE 289

2-(4-{2-Hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-isophthalonitrile

EXAMPLE 290

2-(4-{3-[5-Acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-isophthalonitrile

20

EXAMPLE 291

1-[4-(2-Chloro-6-nitro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol

25

EXAMPLE 292

1-[4-(2-Amino-6-chloro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol

30

EXAMPLE 293

3-Chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzoic acid methyl ester

5

EXAMPLE 294

3-Chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-N-methyl-benzamide

10

EXAMPLE 295

[3-Chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-phenyl]-morpholin-4-yl-methanone

15

EXAMPLE 296

1-[4-(2-Chloro-6-morpholin-4-ylmethyl-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol

20

EXAMPLE 297

3-Chloro-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-N-pyridin-4-ylmethyl-benzamide

25

EXAMPLE 299

2-(4-{2-Hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-3-nitro-benzoic acid methyl ester

30

EXAMPLE 300

2-(4-{3-[5-Acetyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-hydroxy-propyl}-piperazin-1-yl)-3-nitro-benzoic acid methyl ester

5

EXAMPLE 301

3-Acetylamino-2-(4-{2-hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-benzoic acid methyl ester

10

EXAMPLE 302

2-(4-{2-Hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-3-methanesulfonylamino-benzoic acid methyl ester

15

EXAMPLE 303

2-(4-{2-Hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-3-nitro-benzamide

20

EXAMPLE 304

2-(4-{2-Hydroxy-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propyl}-piperazin-1-yl)-3-(3-methyl-ureido)-benzoic acid methyl ester

25

EXAMPLE 305

1-[4-(2,6-Dinitro-phenyl)-piperazin-1-yl]-3-[5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-propan-2-ol

30

EXAMPLE 306

1-[1-{3-[4-(2,6-Dinitro-phenyl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 307

1-[1-{3-[4-(3,5-Dichloro-pyridin-4-yl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

5 EXAMPLE 308

1-{3-[4-(3,5-Dichloro-pyridin-4-yl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

EXAMPLE 309

10 1-[1-{3-[4-(3,5-Dichloro-pyridin-4-yl)-piperazin-1-yl]-2-hydroxy-propyl}-3-(4-trifluoromethylsulfanyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 310

15 1-[1-{3-[4-(3,5-Dichloro-pyridin-4-yl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 311

20 2-(4-{3-[5-Acetyl-3-(4-bromo-phenyl)-4,5,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-1-yl]-2-azido-propyl}-piperazin-1-yl)-benzonitrile

EXAMPLE 312

25 1-[1-{2-Hydroxy-3-[4-(6-nitro-benzothiazol-2-yl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

EXAMPLE 313

1-[1-{2-Hydroxy-3-[4-(6-methoxy-benzothiazol-2-yl)-piperazin-1-yl]-propyl}-3-(4-trifluoromethyl-phenyl)-1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridin-5-yl]-ethanone

30 EXAMPLE 314

1-{3-[4-(1,1-Dioxo-1H-1λ⁶-benzo[d]isothiazol-3-yl)-piperazin-1-yl]-propyl}-5-methanesulfonyl-3-(4-trifluoromethyl-phenyl)-4,5,6,7-tetrahydro-1H-pyrazolo[4,3-c]pyridine

EXAMPLE 315

1-[3-(4-Benzo[d]isothiazol-3-yl-piperazin-1-yl)-propyl]-3-(4-bromo-phenyl)-
1,4,6,7-tetrahydro-pyrazolo[4,3-c]pyridine-5-carboxylic acid amide

5

Table 3

EXAMPLE	IC ₅₀ (μM)
103	2.1
109	1.2
114	1.1
120	12
121	4.3
123	1.9
126	7.4
131	2.2
135	1.4
137	0.98
139	0.71
140	0.53
141	1.4
143	0.35
148	0.63
149	0.86
150	1.8
156	2.9
159	6.2
164	1.9
167	4.9
170	4.5
174	1.4
176	1.6
181	1.7
185	8.4
190	0.26
192	0.79

EXAMPLE (Table 3, cont'd)	IC ₅₀ (μM) (Table 3, cont'd)
193	0.33
195	3.3
196	1.9
199	1.6
205	0.95
208	1.5
211	0.16
214	1.9
216	1.3
219	10
221	0.82
223	0.23
224	0.14
228	2.7
230	0.81
237	1.5
238	3.6
245	0.44
247	4.9
249	0.47
251	4.6
255	0.40
258	0.39
260	0.24
262	0.29
266	0.19
267	0.22
270	1.8
272	0.15

EXAMPLE (Table 3, cont'd)	IC ₅₀ (μM) (Table 3, cont'd)
277	2.8
278	0.19
281	2.5
283	0.08
285	0.04
287	0.20
289	0.15
293	0.14
296	0.48
302	0.07
306	0.34
310	0.40
311	2.1
312	0.77

F. Other Embodiments

The features and advantages of the invention are apparent to one of ordinary skill in the art. Based on this disclosure, including the summary,
5 detailed description, background, examples, and claims, one of ordinary skill in the art will be able to make modifications and adaptations to various conditions and usages. These other embodiments are also within the scope of the invention.

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What is claimed is:

11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100 101 102 103 104 105 106 107 108 109 110 111 112 113 114 115 116 117 118 119 120 121 122 123 124 125 126 127 128 129 130 131 132 133 134 135 136 137 138 139 140 141 142 143 144 145 146 147 148 149 150 151 152 153 154 155 156 157 158 159 160 161 162 163 164 165 166 167 168 169 170 171 172 173 174 175 176 177 178 179 180 181 182 183 184 185 186 187 188 189 190 191 192 193 194 195 196 197 198 199 200 201 202 203 204 205 206 207 208 209 210 211 212 213 214 215 216 217 218 219 220 221 222 223 224 225 226 227 228 229 230 231 232 233 234 235 236 237 238 239 240 241 242 243 244 245 246 247 248 249 250 251 252 253 254 255 256 257 258 259 260 261 262 263 264 265 266 267 268 269 270 271 272 273 274 275 276 277 278 279 280 281 282 283 284 285 286 287 288 289 290 291 292 293 294 295 296 297 298 299 300 301 302 303 304 305 306 307 308 309 310 311 312 313 314 315 316 317 318 319 320 321 322 323 324 325 326 327 328 329 330 331 332 333 334 335 336 337 338 339 340 341 342 343 344 345 346 347 348 349 350 351 352 353 354 355 356 357 358 359 360 361 362 363 364 365 366 367 368 369 370 371 372 373 374 375 376 377 378 379 380 381 382 383 384 385 386 387 388 389 390 391 392 393 394 395 396 397 398 399 400 401 402 403 404 405 406 407 408 409 410 411 412 413 414 415 416 417 418 419 420 421 422 423 424 425 426 427 428 429 430 431 432 433 434 435 436 437 438 439 440 441 442 443 444 445 446 447 448 449 450 451 452 453 454 455 456 457 458 459 460 461 462 463 464 465 466 467 468 469 470 471 472 473 474 475 476 477 478 479 480 481 482 483 484 485 486 487 488 489 490 491 492 493 494 495 496 497 498 499 500 501 502 503 504 505 506 507 508 509 510 511 512 513 514 515 516 517 518 519 520 521 522 523 524 525 526 527 528 529 530 531 532 533 534 535 536 537 538 539 540 541 542 543 544 545 546 547 548 549 550 551 552 553 554 555 556 557 558 559 560 561 562 563 564 565 566 567 568 569 570 571 572 573 574 575 576 577 578 579 580 581 582 583 584 585 586 587 588 589 590 591 592 593 594 595 596 597 598 599 600 601 602 603 604 605 606 607 608 609 610 611 612 613 614 615 616 617 618 619 620 621 622 623 624 625 626 627 628 629 630 631 632 633 634 635 636 637 638 639 640 641 642 643 644 645 646 647 648 649 650 651 652 653 654 655 656 657 658 659 660 661 662 663 664 665 666 667 668 669 670 671 672 673 674 675 676 677 678 679 680 681 682 683 684 685 686 687 688 689 690 691 692 693 694 695 696 697 698 699 700 701 702 703 704 705 706 707 708 709 710 711 712 713 714 715 716 717 718 719 720 721 722 723 724 725 726 727 728 729 730 731 732 733 734 735 736 737 738 739 740 741 742 743 744 745 746 747 748 749 750 751 752 753 754 755 756 757 758 759 760 761 762 763 764 765 766 767 768 769 770 771 772 773 774 775 776 777 778 779 780 781 782 783 784 785 786 787 788 789 790 791 792 793 794 795 796 797 798 799 800 801 802 803 804 805 806 807 808 809 810 811 812 813 814 815 816 817 818 819 820 821 822 823 824 825 826 827 828 829 830 831 832 833 834 835 836 837 838 839 840 841 842 843 844 845 846 847 848 849 850 851 852 853 854 855 856 857 858 859 860 861 862 863 864 865 866 867 868 869 870 871 872 873 874 875 876 877 878 879 880 881 882 883 884 885 886 887 888 889 890 891 892 893 894 895 896 897 898 899 900 901 902 903 904 905 906 907 908 909 910 911 912 913 914 915 916 917 918 919 920 921 922 923 924 925 926 927 928 929 930 931 932 933 934 935 936 937 938 939 940 941 942 943 944 945 946 947 948 949 950 951 952 953 954 955 956 957 958 959 960 961 962 963 964 965 966 967 968 969 970 971 972 973 974 975 976 977 978 979 980 981 982 983 984 985 986 987 988 989 990 991 992 993 994 995 996 997 998 999 1000